

TITLE OF THE INVENTION

LINEAR γ -CARBOXYGLUTAMATE RICH CONOTOXINS

CROSS-REFERENCE TO RELATED APPLICATION

5 [0001] The present application is related to and claims priority under 35 USC §119(e) to U.S. provisional patent application Serial No. 60/273,639 filed 7 March 2001, incorporated herein by reference.

10 [0002] This invention was made with Government support under Grant No. PO1 GM48677 awarded by the National Institute of General Medical Sciences, National Institutes of Health, Bethesda, Maryland. The United States Government has certain rights in the invention.

BACKGROUND OF THE INVENTION

15 [0003] The invention relates to linear γ -carboxyglutamate rich conotoxins, derivatives or pharmaceutically acceptable salts thereof, and uses thereof, including the treatment of neurologic and psychiatric disorders, such as anticonvulsant agents, as neuroprotective agents or for the management of pain. The invention further relates to nucleic acid sequences encoding the conopeptides and encoding propeptides, as well as the propeptides.

20 [0004] The publications and other materials used herein to illuminate the background of the invention, and in particular, cases to provide additional details respecting the practice, are incorporated by reference, and for convenience are referenced in the following text by author and date and are listed alphabetically by author in the appended bibliography.

25 [0005] *Conus* is a genus of predatory marine gastropods (snails) which envenomate their prey. Venomous cone snails use a highly developed projectile apparatus to deliver their cocktail of toxic conotoxins into their prey. In fish-eating species such as *Conus magus* the cone detects the presence of the fish using chemosensors in its siphon and when close enough extends its proboscis and fires a hollow harpoon-like tooth containing venom into the fish. The venom immobilizes the fish and enables the cone snail to wind it into its mouth via an attached filament. For general information on *Conus* and their venom see the website address 30 <http://grimwade.biochem.unimelb.edu.au/cone/referenc.html>. Prey capture is accomplished through a sophisticated arsenal of peptides which target specific ion channel and receptor subtypes. Each *Conus* species venom appears to contain a unique set of 50-200 peptides. The composition of the venom differs greatly between species and between individual snails within each species, each optimally evolved to paralyse it's prey. The active components of the venom

are small peptides toxins, typically 12-30 amino acid residues in length and are typically highly constrained peptides due to their high density of disulphide bonds.

[0006] The venoms consist of a large number of different peptide components that when separated exhibit a range of biological activities: when injected into mice they elicit a range of physiological responses from shaking to depression. The paralytic components of the venom that have been the focus of recent investigation are the α -, ω - and μ -conotoxins. All of these conotoxins act by preventing neuronal communication, but each targets a different aspect of the process to achieve this. The α -conotoxins target nicotinic ligand gated channels, the μ -conotoxins target the voltage-gated sodium channels and the ω -conotoxins target the voltage-gated calcium channels (Olivera et al., 1985; Olivera et al., 1990). For example a linkage has been established between α -, α A- & ϕ -conotoxins and the nicotinic ligand-gated ion channel; ω -conotoxins and the voltage-gated calcium channel; μ -conotoxins and the voltage-gated sodium channel; δ -conotoxins and the voltage-gated sodium channel; κ -conotoxins and the voltage-gated potassium channel; conantokins and the ligand-gated glutamate (NMDA) channel.

[0007] However, the structure and function of only a small minority of these peptides have been determined to date. For peptides where function has been determined, three classes of targets have been elucidated: voltage-gated ion channels; ligand-gated ion channels, and G-protein-linked receptors.

[0008] *Conus* peptides which target voltage-gated ion channels include those that delay the inactivation of sodium channels, as well as blockers specific for sodium channels, calcium channels and potassium channels. Peptides that target ligand-gated ion channels include antagonists of NMDA and serotonin receptors, as well as competitive and noncompetitive nicotinic receptor antagonists. Peptides which act on G-protein receptors include neuropeptides and vasopressin receptor agonists. The unprecedented pharmaceutical selectivity of conotoxins is at least in part defined by a specific disulfide bond frameworks combined with hypervariable amino acids within disulfide loops (for a review see McIntosh et al., 1998).

[0009] The conantokins are structurally unique. In contrast to the well characterized conotoxins from *Conus* venoms, most conantokins do not contain disulfide bonds. However, they contain 4-5 residues of the unusual modified amino acid γ -carboxyglutamic acid. The occurrence of this modified amino acid, which is derived post-translationally from glutamate in a vitamin K-dependent reaction, was unprecedented in a neuropeptide. It has been established that the conantokins have N-methyl-D-aspartate (NMDA) antagonist activity, and consequently

target the NMDA receptor. The conantokins reduce glutamate (or NMDA) mediated increases in intracellular Ca^{2+} and cGMP without affecting kainate-mediated events (Chandler et al., 1993). Although these peptides have actions through polyamine responses of the NMDA receptors, the neurochemical profile of these polypeptides is distinct from previously described 5 noncompetitive NMDA antagonists (Skolnick et al., 1992).

[0010] Ischemic damage to the central nervous system (CNS) may result from either global or focal ischemic conditions. Global ischemia occurs under conditions in which blood flow to the entire brain ceases for a period of time, such as may result from cardiac arrest. Focal ischemia occurs under conditions in which a portion of the brain is deprived of its normal blood supply, such as may result from thromboembolic occlusion of a cerebral vessel, traumatic head or spinal cord injury, edema or brain or spinal cord tumors. Both global and focal ischemic conditions have the potential for widespread neuronal damage, even if the global ischemic condition is transient or the focal condition affects a very limited area.

[0011] Epilepsy is a recurrent paroxysmal disorder of cerebral function characterized by sudden brief attacks of altered consciousness, motor activity, sensory phenomena or inappropriate behavior caused by abnormal excessive discharge of cerebral neurons. Convulsive seizures, the most common form of attacks, begin with loss of consciousness and motor control, and tonic or clonic jerking of all extremities but any recurrent seizure pattern may be termed epilepsy. The term primary or idiopathic epilepsy denotes those cases where no cause for the 20 seizures can be identified. Secondary or symptomatic epilepsy designates the disorder when it is associated with such factors as trauma, neoplasm, infection, developmental abnormalities, cerebrovascular disease, or various metabolic conditions. Epileptic seizures are classified as partial seizures (focal, local seizures) or generalized seizures (convulsive or nonconvulsive).

25 Classes of partial seizures include simple partial seizures, complex partial seizures and partial seizures secondarily generalized. Classes of generalized seizures include absence seizures, atypical absence seizures, myoclonic seizures, clonic seizures, tonic seizures, tonic-clonic seizures (*grand mal*) and atonic seizures. Therapeutics having anticonvulsant properties are used in the treatment of seizures. Most therapeutics used to abolish or attenuate seizures act at least through effects that reduce the spread of excitation from seizure foci and prevent 30 detonation and disruption of function of normal aggregates of neurons. Traditional anticonvulsants that have been utilized include phenytoin, phenobarbital, primidone, carbamazepine, ethosuximide, clonazepam and valproate. Several novel and chemically diverse anticonvulsant medications recently have been approved for marketing, including lamotrigine,

ferbamate, gabapentin and topiramate. For further details of seizures and their therapy, see Rall & Schleifer (1985) and *The Merck Manual* (1992).

[0012] (S)-Glutamic acid (Glu), which is the main excitatory neurotransmitter in the CNS, and other excitatory amino acids (EAA) operate through four different classes of receptors. In addition to the three heterogeneous classes of ionotropic EAA receptors (iGluRs), named M-methyl-D-aspartate (NMDA), (RS)-2-amino-3-(hydroxy-5-methyl-4-isoxazolyl)-propionic acid (AMPA) and Kainate (KA) receptors, a heterogeneous class of G-protein coupled EAA receptors (mGluRs) has been shown to have important functions in neuronal signalling processes. It is now generally agreed that iGluRs as well as mGluRs play important roles in the healthy as well as the diseased CNS, and that all subtypes of these receptors are potential targets for therapeutic intervention in a number of diseases. For a review, see Brauner-Osborne et al. (2000).

[0013] The cloning of the different subunits of the iGluRs and of the eight subtypes of mGluRs represents a major breakthrough. Whereas at present six NMDA receptor subunits (NR1, NR2A-NR2D, and NR3A) have been cloned and characterised in regards to primary structure, four AMPA subunits (iGluR1-4) have similarly been characterized, and so far 5 subunits building blocks for KA-preferred receptors (iGluR5-7, KA1, and KA2) have been identified. Most if not all physiological iGluRs have heterotetra- or pentameric structures, but the number of functional NMDA, AMPA, and KA receptors in the CNS is not known. At present 8 subtypes of the 7TM mGluRs have been characterized, but there is evidence to suggest that further subtypes of mGluRs may be identified. The structurally unique linear conantokin peptides disclosed in this patent represent a series of ligands capable of activating, blocking or allosterically modulating both iGluRs and mGluRs – they represent essential pharmacological tools and potential therapeutics for treatment brain injury, stroke, Huntingdon's disease, Parkinson's disease, Alzheimers disease, ALS, Epilepsy, Schizophrenia, pain, anxiety, AIDS related dementia, spinal injury amongst other chronic and acute diseases and conditions.

[0014] For example, the NMDA receptor is involved in a broad spectrum of CNS disorders. For example, during brain ischemia caused by stroke or traumatic injury, excessive amounts of the excitatory amino acid glutamate are released from damaged or oxygen deprived neurons. This excess glutamate binds the NMDA receptor which opens the ligand-gated ion channel thereby allowing Ca^{2+} influx producing a high level of intracellular Ca^{2+} , which activates biochemical cascades resulting in protein, DNA and membrane degradation leading to cell death. This phenomenon, known as excitotoxicity, is also thought to be responsible for the

neurological damage associated with other disorders ranging from hypoglycemia and cardiac arrest to epilepsy. In addition, there are reports indicating similar involvement in the chronic neurodegeneration of Huntington's, Parkinson's and Alzheimer's diseases.

[0015] Parkinson's disease is a progressive, neurodegenerative disorder. The etiology of the disorder is unknown in most cases, but has been hypothesized to involve oxidative stress. The underlying neuropathology in Parkinsonian patients is an extensive degenerations of the pigmented dopamine neurons in the substantia nigra. These neurons normally innervate the caudate and putamen nuclei. Their degeneration results in a marked loss of the neurotransmitter dopamine in the caudate and putamen nuclei. This loss of dopamine and its regulation of neurons in the caudate-putamen leads to the bradykinesia, rigidity, and tremor that are the hallmarks of Parkinson's disease. An animal model has been developed for Parkinson's disease (Zigmond et al., 1987) and has been used to test agents for anti-Parkinsonian activity (Ungerstedt et al., 1973).

[0016] The dopamine precursor, L-Dopa, is the current therapy of choice in treating the symptoms of Parkinson's disease. However, significant side effects develop with continued use of this drug and with disease progression, making the development of novel therapies important. Recently, antagonists of the NMDA subtype of glutamate receptor have been proposed as potential anti-Parkinsonian agents. (Borman, 1989; Greenamyre and O'Brien, 1991; Olney et al., 1987). In addition, antagonists of NMDA receptors potentiate the behavioral effects of L-Dopa and D1 dopamine receptor stimulation in animal models of Parkinson's disease. (Starr, 1995). These data suggest that NMDA receptor antagonists may be useful adjuncts to L-Dopa therapy in Parkinson's disease by decreasing the amount of L-Dopa required and thereby reducing undesirable side effects. In addition, antagonists of NMDA receptors have been shown to attenuate free radical mediated neuronal death. Thus, NMDA receptor antagonists may also prevent further degeneration of dopamine neurons in addition to providing symptomatic relief. Finally, NMDA receptor antagonists have been shown to potentiate the contralateral rotations induced by L-Dopa or D1 dopamine receptor antagonists in the animal model.

[0017] Pain, and particularly, persistent pain, is a complex phenomenon involving many interacting components. Numerous studies, however, have demonstrated a role for NMDA receptors in mediating persistent pain, and further that NMDA antagonists are effective in animal models of persistent pain. See for example, PCT published application WO 98/03189.

[0018] Neuropsychiatric involvement of the NMDA receptor has also been recognized. Blockage of the NMDA receptor Ca²⁺ channel by the animal anesthetic phencyclidine produces

a psychotic state in humans similar to schizophrenia (Johnson et al., 1990). Further, NMDA receptors have also been implicated in certain types of spatial learning (Bliss et al., 1993). In addition, numerous studies have demonstrated a role for NMDA receptors in phenomena associated with addiction to and compulsive use of drugs or ethanol. Furthermore, antagonists 5 of NMDA receptors may be useful for treating addiction-related phenomena such as tolerance, sensitization, physical dependence and craving (for review see, Popik et al., 1995; Spanagel and Zieglgansberger, 1997; Trujillo and Akil, 1995).

[0019] There are several lines of evidence which suggest that NMDA antagonists may be useful in the treatment of HIV infection. First, the levels of the neurotoxin and NMDA agonist quinolinic acid are elevated in the cerebrospinal fluid of HIV-positive subjects (Heyes et al., 1989) and in murine retrovirus-induced immunodeficiency syndrome (Sei et al., 1996). Second, the envelope glycoprotein of HIV-1 alters NMDA receptor function (Sweetnam et al., 1993). Thirdly, NMDA antagonists can reduce the effects and neurotoxicity of GP-120 (Muller et al., 1996; Raber et al., 1996; Nishida et al., 1996). Fourth, GP-120 and glutamate act synergistically to produce toxicity *in vitro* (Lipton et al., 1991). And finally, memantine, an NMDA antagonist, protects against HIV infection in glial cells *in vitro* (Rytik et al., 1991). For 10 a review of the use of NMDA antagonists in treating HIV infection, see Lipton (1994; 1996). For 15

[0020] PCT published application WO 98/03189 has shown that the class of conopeptides termed conantokins are useful for treating each of the previously discussed 20 disorders as well as several others, including mood disorders, urinary incontinence, dystonia and sleep disorders among others. U.S. Patent No. 5,844,077 also discloses the use of conantokins for inducing analgesia and for neuroprotection.

[0021] It is desired to identify additional compounds which are useful as anticonvulsant, 25 neuroprotective, neuropsychiatric or analgesic agents.

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SUMMARY OF THE INVENTION

[0022] The present invention is directed to linear γ -carboxyglutamate rich conotoxins, derivatives or pharmaceutically acceptable salts thereof, and uses thereof, including the treatment of neurologic and psychiatric disorders, such as anticonvulsant agents, as 30 neuroprotective agents or for the management of pain. The invention is further directed to nucleic acid sequences encoding the conopeptides and encoding propeptides, as well as the propeptides.

[0023] More specifically, the present invention is directed to linear γ -carboxyglutamate rich conotoxins, having the amino acid sequences:

Conotoxin-Af6: X₆GQDDSX₁X₁X₁DSQX₂VMX₂HGQRERR[^] (SEQ ID NO:1)

Conotoxin-Bt1: GGX₁X₁VRX₁SAX₁TLHX₁LTX₅[^] (SEQ ID NO:2)

5 Conotoxin-Bt2: GGX₁X₁VRX₁SAX₁TLHX₁ITX₅[^] (SEQ ID NO:3)

Conotoxin-Bt3: DGX₁X₁VRX₁AAX₁TLNX₁LTX₅[^] (SEQ ID NO:4)

Conotoxin-Bt4: GYX₁DDRX₁IAX₁TVRX₁LX₁X₁A# (SEQ ID NO:5)

Conotoxin-Bt5: GGGX₁VRX₁SAX₁TLHX₁ITX₅[^] (SEQ ID NO:6)

Conotoxin-Bu1: NX₅X₁TX₃IX₁IVX₁ISRX₁LX₁X₁I# (SEQ ID NO:7)

Conotoxin-Bu2: NX₅X₁TX₃X₃NLX₁LVX₁ISRX₁LX₁X₁I# (SEQ ID NO:8)

Conotoxin-C1: SDX₁X₁LLRX₁DVX₁TVLX₁LX₁RN# (SEQ ID NO:9)

Conotoxin-C2: GDX₁X₁LLRX₁DVX₁TVLX₁LX₁RD# (SEQ ID NO:10)

Conotoxin-C3: SDX₁X₁LLRX₁DVX₁TVLX₁PX₁RN# (SEQ ID NO:11)

Conotoxin-C4: IX₁X₁GLIX₁DLX₁TARX₁RDS# (SEQ ID NO:12)

Conotoxin-C5: IX₁X₁GLIX₁DLX₁AARX₁RDS# (SEQ ID NO:13)

Conotoxin-C6: GX₁X₅X₁VGSIX₅X₁AVRQQX₁CIRNNNNRX₅X₄CX₅X₂[^] (SEQ ID NO:14)

Conotoxin-Di1: TITAX₁X₁AX₁RTSX₁RMSSM# (SEQ ID NO:15)

Conotoxin-Di2: X₆X₁TX₅TX₅X₁X₁VX₁RHTX₁RLKSM# (SEQ ID NO:16)

20 Conotoxin-Ep1: GGKDIVX₁TITX₁LX₁X₂I# (SEQ ID NO:17)

Conotoxin-Fi1: GX₁X₁X₁VAX₁MAAX₁IARX₁NQAN# (SEQ ID NO:18)

Conotoxin-Fi2: SX₃X₁QARX₁VQX₁AVNX₁LX₂X₁R# (SEQ ID NO:19)

Conotoxin-Fi2a: SX₃X₁QARX₁VQX₁AVNX₁LX₂X₁RGX₂X₂IIMLGVX₅R-DTRQF[^] (SEQ ID NO:20)

25 Conotoxin-Fi3: D X₃X₁DDRX₁IAX₁TVRX₁LX₁X₁I# (SEQ ID NO:21)

Conotoxin-Fi4: GNTAX₁X₁VRX₁AAX₁TLHX₁LSL[^] (SEQ ID NO:22)

Conotoxin-Fi5: GSISM GFX₁HRRX₁IAX₁LVRX₁LAX₁I# (SEQ ID NO:23)

Conotoxin-L1: GX₁X₁X₁VAX₁MAAX₁IARX₁NAAN# (SEQ ID NO:24)

Conotoxin-L2: GX₂X₁DRX₁IVX₁TVRX₁LX₁X₁I# (SEQ ID NO:25)

30 Conotoxin-L3: GX₁X₁X₁VAX₂MAAX₁LTRX₁X₁AVX₂# (SEQ ID NO:26)

Conotoxin-P1: GX₁X₁X₁HSX₂X₃QX₁CLRX₁VRVNX₂VQQX₁C[^] (SEQ ID NO:27)

Conotoxin-P2:	$\text{GX}_1\text{X}_1\text{X}_1\text{HSX}_2\text{X}_3\text{QX}_1\text{CLRX}_1\text{VRVNNVQQX}_1\text{C}^{\wedge}$	(SEQ	ID
NO:28)			
Conotoxin-P3:	$\text{GX}_1\text{X}_1\text{X}_1\text{HSX}_2\text{X}_3\text{QX}_1\text{CLRX}_1\text{IRVNX}_2\text{VQQX}_1\text{C}^{\wedge}$	(SEQ	ID
NO:29)			
5 Conotoxin-P4:	$\text{GX}_1\text{AX}_1\text{HX}_3\text{AFQX}_1\text{CLRX}_1\text{INVNX}_2\text{VQQX}_1\text{C}^{\wedge}$	(SEQ ID NO:30)	
Conotoxin-P5:	$\text{GLX}_1\text{X}_1\text{DIX}_1\text{FIX}_1\text{TIX}_1\text{X}_1\text{I}^{\#}$	(SEQ ID NO:31)	
Conotoxin-Sm1:	$\text{ITX}_1\text{TDIX}_1\text{LVMX}_2\text{LX}_1\text{X}_1\text{I}^{\#}$	(SEQ ID NO:32)	

[0024] wherein X_1 is Glu or γ -carboxyglutamic acid (Gla); X_2 is Lys, nor-Lys, N-methyl-Lys, N,N-dimethyl-Lys or N,N,N-trimethyl-Lys; X_3 is Tyr, mono-halo-Tyr, di-halo-Tyr, O-sulpho-Tyr, O-phospho-Tyr or nitro-Tyr; X_4 is Trp (D or L) or halo-Trp (D or L); X_5 is Pro or hydroxy-Pro; and X_6 is Gln or pyroglutamate. The halo is preferably chlorine, bromine or iodine, more preferably iodine for Tyr and bromine for Trp. The C-terminus contains a carboxyl or an amide. The preferred C-terminus is shown herein in Tables 5 and 6, which shows an alignment of the conopeptides of the present invention.

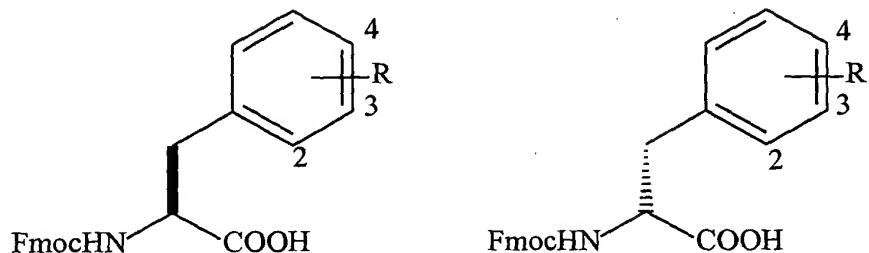
[0025] The present invention is further directed to derivatives or pharmaceutically acceptable salts of the linear γ -carboxyglutamate rich conotoxins or their derivatives. Examples of derivatives include peptides in which the γ -carboxyglutamic acid at the X_1 residues of the peptides of the present invention other than those residues corresponding to residues 3 and 4 of conatntokin G, such as shown by the alignment set forth herein in Table 5 by X, is replaced by any other amino acids such that their NMDA antagonist activity is not adversely affected. Examples of such replacements include, but are not limited to Ser, Ala, Glu and Tyr. Other derivatives are produced by modification of the amino acids within the peptide structure. Modified amino acids include those which are described in Roberts et al. (1983). Other derivatives include peptides in which one or more residues have been deleted. It has been discovered that one to five of the C-terminal amino acid residues can be deleted without loss of activity. Substitutions of one amino acid for another can be made at one or more additional sites within the above peptide, and may be made to modulate one or more of the properties of the peptides. Substitutions of this kind are preferably conservative, i.e., one amino acid is replaced with one of similar shape and charge. Conservative substitutions are well known in the art and include, for example: alanine to glycine, arginine to lysine, asparagine to glutamine or histidine, glycine to proline, leucine to valine or isoleucine, serine to threonine, phenylalanine to tyrosine, and the like.

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[0026] These derivatives also include peptides in which the Arg residues may be substituted by Lys, ornithine, homoarginine, nor-Lys, N-methyl-Lys, N,N-dimethyl-Lys, N,N,N-trimethyl-Lys or any synthetic basic amino acid; the Lys residues may be substituted by Arg, ornithine, homoarginine, nor-Lys, or any synthetic basic amino acid; the Tyr residues may be substituted with meta-Tyr, ortho-Tyr, nor-Tyr, mono-halo-Tyr, di-halo-Tyr, O-sulpho-Tyr, O-phospho-Tyr, nitro-Tyr or any synthetic hydroxy containing amino acid; the Ser residues may be substituted with Thr or any synthetic hydroxylated amino acid; the Thr residues may be substituted with Ser or any synthetic hydroxylated amino acid; the Phe residues may be substituted with any synthetic aromatic amino acid; the Trp residues may be substituted with Trp (D), neo-Trp, halo-Trp (D or L) or any aromatic synthetic amino acid; and the Asn, Ser, Thr or Hyp residues may be glycosylated. The halogen may be iodo, chloro, fluoro or bromo; preferably iodo for halogen substituted-Tyr and bromo for halogen-substituted Trp. The Tyr residues may also be substituted with the 3-hydroxyl or 2-hydroxyl isomers (meta-Tyr or ortho-Tyr, respectively) and corresponding O-sulpho- and O-phospho-derivatives. The acidic amino acid residues may be substituted with any synthetic acidic amino acid, e.g., tetrazolyl derivatives of Gly and Ala. The Met residues may be substituted with norleucine (Nle). The aliphatic amino acids may be substituted by synthetic derivatives bearing non-natural aliphatic branched or linear side chains C_nH_{2n+2} up to and including n=8.

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[0027] Examples of synthetic aromatic amino acid include, but are not limited to, nitro-Phe, 4-substituted-Phe wherein the substituent is C_1 - C_3 alkyl, carboxyl, hydroxymethyl, sulphomethyl, halo, phenyl, -CHO, -CN, - SO_3H and -NHAc. Examples of synthetic hydroxy containing amino acid, include, but are not limited to, such as 4-hydroxymethyl-Phe, 4-hydroxyphenyl-Gly, 2,6-dimethyl-Tyr and 5-amino-Tyr. Examples of synthetic basic amino acids include, but are not limited to, N-1-(2-pyrazolinyl)-Arg, 2-(4-piperinyl)-Gly, 2-(4-piperinyl)-Ala, 2-[3-(2S)pyrrolinyl]-Gly and 2-[3-(2S)pyrrolinyl]-Ala. These and other synthetic basic amino acids, synthetic hydroxy containing amino acids or synthetic aromatic amino acids are described in Building Block Index, Version 3.0 (1999 Catalog, pages 4-47 for hydroxy containing amino acids and aromatic amino acids and pages 66-87 for basic amino acids; see also <http://www.amino-acids.com>), incorporated herein by reference, by and available from RSP Amino Acid Analogues, Inc., Worcester, MA. Examples of synthetic acid amino acids include those derivatives bearing acidic functionality, including carboxyl, phosphate, sulfonate and synthetic tetrazolyl derivatives such as described by Ornstein et al. (1993) and in

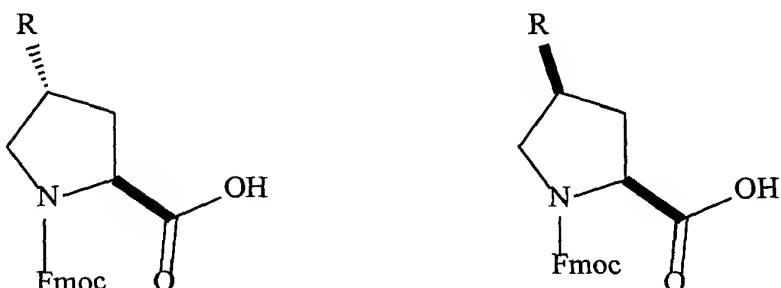
U.S. Patent No. 5,331,001, each incorporated herein by reference, and such as shown in the following schemes 1-3.

Tetrazole Compounds



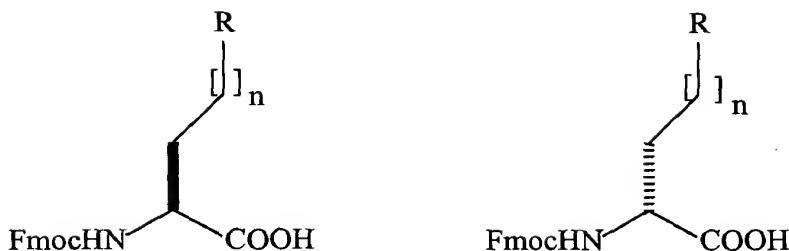
R=COOH, tetrazole, CH_2COOH , 4-NHSO₂CH₃, 4-NHSO₂Phenyl,
 4-CH₂SO₃H, SO₃H, 4-CH₂PO₃H₂, CH₂CH₂COOH, OCH₂Tetrazole,
 CH₂STetrazole, HNTetrazole, CONHSO₂R₁ where R₁ is CH₃ or Phenyl
 SO₂-Tetrazole, CH₂CH₂SO₃H, 1,2,4-tetrazole, 3-isoxazolone,
 amidotetrazole, CH₂CH₂PO₃H₂

Scheme 1



R = COOH, tetrazole, CH_2COOH , CH_2 tetrazole

Scheme 2



R = COOH, tetazole, CH₂COOH, 4-NHSO₂CH₃, 4-NHSO₂Phenyl, 4-CH₂SO₃H, SO₃H, 4-CH₂PO₃H₂, CH₂CH₂COOH, OCH₂Tetrazole, CH₂STetrazole, HNTetrazole, CONHSO₂R₁ where R₁ is CH₃ or Phenyl SO₂-Tetrazole, CH₂CH₂SO₃H, 1,2,4-tetrazole, 3-isoxazolone, amidotetrazole, CH₂CH₂PO₃H₂ n = 0, 1, 2, or 3

Scheme 3

PCT/US2007/035007

[0028] Optionally, in the linear γ -carboxyglutamate rich conotoxins of the present invention, the Asn residues may be modified to contain an N-glycan and the Ser, Thr and Hyp residues may be modified to contain an O-glycan (e.g., g-N, g-S, g-T and g-Hyp). In accordance with the present invention, a glycan shall mean any N-, S- or O-linked mono-, di-, tri-, poly- or oligosaccharide that can be attached to any hydroxy, amino or thiol group of natural or modified amino acids by synthetic or enzymatic methodologies known in the art. The monosaccharides making up the glycan can include D-allose, D-altrose, D-glucose, D-mannose, D-gulose, D-10 idose, D-galactose, D-talose, D-galactosamine, D-glucosamine, D-N-acetyl-glucosamine (GlcNAc), D-N-acetyl-galactosamine (GalNAc), D-fucose or D-arabinose. These saccharides may be structurally modified, e.g., with one or more O-sulfate, O-phosphate, O-acetyl or acidic groups, such as sialic acid, including combinations thereof. The glycan may also include similar polyhydroxy groups, such as D-penicillamine 2,5 and halogenated derivatives thereof or polypropylene glycol derivatives. The glycosidic linkage is beta and 1-4 or 1-3, preferably 1-3. The linkage between the glycan and the amino acid may be alpha or beta, preferably alpha and is 1-.

[0029] Core O-glycans have been described by Van de Steen et al. (1998), incorporated herein by reference. Mucin type O-linked oligosaccharides are attached to Ser or Thr (or other 20 hydroxylated residues of the present peptides) by a GalNAc residue. The monosaccharide building blocks and the linkage attached to this first GalNAc residue define the "core glycans," of which eight have been identified. The type of glycosidic linkage (orientation and

connectivities) are defined for each core glycan. Suitable glycans and glycan analogs are described further in U.S. Serial No. 09/420,797 filed 19 October 1999 and in PCT Application No. PCT/US99/24380 filed 19 October 1999 (PCT Published Application No. WO 00/23092), each incorporated herein by reference. A preferred glycan is Gal(β1→3)GalNAc(α1→).

5 [0030] More specifically, the present invention is also directed to nucleic acids which encode linear γ -carboxyglutamate rich conotoxins of the present invention or which encodes precursor peptides for these conotoxins, as well as the precursor peptide. The nucleic acid sequences encoding the precursor peptides of other conopeptides of the present invention are set forth in Table 4.

10 [0031] The present invention is further directed to uses of these peptides or nucleic acids as described herein, including the treatment of neurologic and psychiatric disorders, such as anticonvulsant agents, as neuroprotective agents or for the management of pain.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

15 [0032] The present invention is directed to linear γ -carboxyglutamate rich conotoxins, derivatives or pharmaceutically acceptable salts thereof. The present invention is further directed to the use of this peptide, derivatives thereof and pharmaceutically acceptable salts thereof for the treatment of neurologic and psychiatric disorders, such as anticonvulsant agents, as neuroprotective agents or for the management of pain, e.g. as analgesic agents. Neurologic disorders and psychiatric disorders as used herein are intended to include such disorders as grouped together in *The Merck Manual of Diagnosis and Therapy*, inclusive of the disorders discussed in PCT published application WO 98/03189, incorporated herein by reference. The invention is further directed to nucleic acid sequences encoding the conopeptides and encoding propeptides, as well as the propeptides.

20 [0033] More specifically, the present invention is directed to the use of these compounds for the treatment and alleviation of epilepsy and as a general anticonvulsant agent. The present invention is also directed to the use of these compounds for reducing neurotoxic injury associated with conditions of hypoxia, anoxia or ischemia which typically follows stroke, cerebrovascular accident, brain or spinal cord trauma, myocardial infarct, physical trauma, drowning, suffocation, perinatal asphyxia, or hypoglycemic events. The present invention is further directed to the use of these compounds for treating neurodegeneration associated with Alzheimer's disease, senile dementia, Amyotrophic Lateral Sclerosis, Multiple Sclerosis, Parkinson's disease, Huntington's disease, Down's Syndrome, Korsakoff's disease,

schizophrenia, AIDS dementia, multi-infarct dementia, Binswanger dementia and neuronal damage associated with uncontrolled seizures. The present invention is also directed to the use of these compounds for treating chemical toxicity, such as addiction, drug craving, alcohol abuse, morphine tolerance, opioid tolerance and barbiturate tolerance. The present invention is further directed to treating psychiatric disorders, such as anxiety, major depression, manic-depressive illness, obsessive-compulsive disorder, schizophrenia and mood disorders (such as bipolar disorder, unipolar depression, dysthymia and seasonal effective disorder). These compounds are also useful for treating ophthalmic disorders. The present invention is also directed to treating additional neurological disorders, such as dystonia (movement disorder), sleep disorder, muscle relaxation and urinary incontinence. In addition, these compounds are useful for memory/cognition enhancement, i.e., treating memory, learning or cognitive deficits. The present invention is also useful in the treatment of HIV infection. Finally, the present invention is directed to the use of these compounds for controlling pain, e.g. as analgesic agents, and the treatment of migraine, acute pain or persistent pain. They can be used prophylactically and also to relieve the symptoms associated with a migraine episode.

[0034] The conopeptides, their derivatives and their salts, have anticonvulsant activity in Frings audiogenic seizure susceptible mice and in syndrome-specific seizure animal models. These peptides also have activity in animal pain models. These peptides further have activity in *in vitro* assays for protection from neurotoxicity. These peptides also have activity in animal models for Parkinson's disease. Thus, the peptides of the present invention are useful as anticonvulsant agents, as neuroprotective agents, as analgesic agents, for managing pain and for treating neurodegenerative disorders. The peptides are administered to patients as described further below.

[0035] These peptides are sufficiently small to be chemically synthesized. General chemical syntheses for preparing the foregoing peptides are described in PCT published application WO 98/03189. The peptides are synthesized by a suitable method, such as by exclusively solid-phase techniques, by partial solid-phase techniques, by fragment condensation or by classical solution couplings. The peptides are also synthesized using an automatic synthesizer. Conopeptides of the present invention can also be obtained by isolation and purification from specific *Conus* species using the technique described in PCT published application WO 98/03189.

[0036] Although the conopeptides of the present invention can be obtained by purification from cone snails, because the amounts of peptide obtainable from individual snails are very

small, the desired substantially pure peptides are best practically obtained in commercially valuable amounts by chemical synthesis using solid-phase strategy. For example, the yield from a single cone snail may be about 10 micrograms or less of peptide. By "substantially pure" is meant that the peptide is present in the substantial absence of other biological molecules of the same type; it is preferably present in an amount of at least about 85% purity and preferably at least about 95% purity.

[0037] The peptides of the present invention can also be produced by recombinant DNA techniques well known in the art. Such techniques are described by Sambrook et al. (1989). The peptides produced in this manner are isolated, reduced if necessary, and oxidized, if necessary, to form the correct disulfide bonds.

[0038] The conopeptides of the present invention have been found to be antagonists of the excitatory amino acid (EAA) receptors, including the ionotropic glutamate (or EAA) receptors (iGluRs, including NMDA receptors, AMPA receptors and KA receptors) and the G-protein coupled glutamate (or EAA) receptors (mGluRs). For example, conopeptide JG001, has been found to be an antagonist of the NMDA receptor subunits and is useful as anticonvulsant agents, as neuroprotective agents, as analgesic agents, for managing pain and for treating neurodegenerative disorders. The conopeptides of the present invention, as well as their derivatives and salts, are particularly useful as such agents for treating neurologic disorders and psychiatric disorders that result from an overstimulation of excitatory amino acid receptors. That is, the invention pertains particularly to disorders in which the pathophysiology involves excessive excitation of nerve cells by excitatory amino acids or agonists of the ionotropic EAA receptors, such as the NMDA receptor(s), AMPA receptor and KA receptor and of the G-protein coupled EAA receptors. Thus, the conopeptides of the present invention are useful for the treatment and alleviation of epilepsy and as general anticonvulsant agents. The use of the conopeptides of the present invention in these conditions includes the administration of a conopeptide in a therapeutically effective amount to patients in need of treatment. The conopeptides of the present invention can be used to treat the seizures, to reduce their effects and to prevent seizures.

[0039] The conopeptides of the present invention are also useful to reduce neurotoxic injury associated with conditions of hypoxia, anoxia or ischemia which typically follows stroke, cerebrovascular accident, brain or spinal chord trauma, myocardial infarct, physical trauma, drownings, suffocation, perinatal asphyxia, or hypoglycemic events. To reduce neurotoxic injury, a conopeptide should be administered in a therapeutically effective amount to the patient

within 24 hours of the onset of the hypoxic, anoxic or ischemic condition in order for conopeptide to effectively minimize the CNS damage which the patient will experience.

[0040] The conopeptides are further useful for the treatment of Alzheimer's disease, senile dementia, Amyotrophic Lateral Sclerosis, Multiple Sclerosis, Parkinson's disease, 5 Huntington's disease, Down's Syndrome, Korsakoff's disease, schizophrenia, AIDS dementia, multi-infarct dementia, Binswanger dementia and neuronal damage associated with uncontrolled seizures. The administration of a conopeptide in a therapeutically effective amount to a patient experiencing such conditions will serve to either prevent the patient from experiencing further neurodegeneration or it will decrease the rate at which neurodegeneration occurs. In addition, the conopeptides can be administered in adjunct with conventional treatment agents to reduce the amount of such agents which need to be used.

[0041] The conopeptides of the present invention are also useful for treating chemical toxicity (such as addiction, morphine tolerance, opiate tolerance, opioid tolerance and barbiturate tolerance), anxiety, major depression, manic-depressive illness, obsessive-compulsive disorder, schizophrenia, mood disorders (such as bipolar disorder, unipolar depression, dysthymia and seasonal effective disorder), dystonia (movement disorder), sleep disorder, muscle relaxation, urinary incontinence, HIV infection and ophthalmic indications. In treating these conditions, a therapeutically effective amount of a conopeptide is administered to a patient to completely treat the condition or to ease the effects of the condition. In addition, the conopeptides are useful for 20 memory/cognition enhancement (treating memory, learning or cognitive deficits), in which case a therapeutically effective amount of a conopeptide is administered to enhance memory or cognition.

[0042] The conopeptides of the present invention are further useful in controlling pain, e.g., as analgesic agents, and the treatment of migraine, acute pain or persistent pain. They can 25 be used prophylactically or to relieve the symptoms associated with a migraine episode, or to treat acute or persistent pain. For these uses, a conopeptide is administered in a therapeutically effective amount to overcome or to ease the pain.

[0043] The anticonvulsant effects of the conopeptide JG001 has been demonstrated in 30 animal models. In rodents, conopeptide JG001 is effective against supramaximal tonic extension seizures produced by maximal electroshock and threshold seizures induced by subcutaneous (s.c.) pentylenetetrazole or picrotoxin. As described in further detail below, conopeptide JG001 was found to have a protective index of 20. Conopeptide JG001 is also effective against focal seizures induced by aluminum hydroxide injection into the pre- and post-

central gyri of rhesus monkeys. Conopeptide JG001, when administered to patients with refractory complex partial seizures, may markedly reduce seizure frequency and severity. Thus, conopeptide JG001 is useful as anticonvulsant agents. Moreover, the clinical utility of conopeptide JG001 as a therapeutic agent for epilepsy may include generalized tonic-clonic and complex partial seizures.

[0044] The neuroprotective effects of conopeptide JG001 is demonstrated in laboratory animal models. In these models, conopeptide JG001 protects against hypoxic damage to the hippocampal slice *in vitro*. In neonate rats, conopeptide JG001 reduces the size of cortical infarcts and amount of hippocampal necrosis following bilateral carotid ligation and hypoxia. Thus, conopeptide JG001 are useful as neuroprotective agents. Whereas other anticonvulsants may exhibit neuroprotectant properties (Aldrete et al., 1979; Abiko et al., 1986; Nehlig et al., 1990), these effects often occurred only at high, clinically achievable doses associated with considerable toxicity (Troupin et al., 1986; Wong et al., 1986). In contrast, conopeptide JG001 exhibits both anticonvulsant and neuroprotectant effects at doses well tolerated by animals and humans.

[0045] The analgesic or anti-pain activity of conopeptide JG001 is demonstrated in animal models of pain and in animal models of persistent pain. In these models, conopeptide JG001 is (a) effective in nerve injury model studies; (b) effective in reducing the tolerance to opiate analgesics after chronic administration and (c) effective in inhibiting activation of NMDA receptors and thereby inhibiting the release of Substance P by small-diameter, primary, sensory pain fibers. Thus, conopeptide JG001 is useful as analgesic agents and anti-pain agents for the treatment of acute and persistent pain. Conopeptide JG001 is also useful for treating addiction, morphine/opiate/opioid tolerance or barbiturate tolerance.

[0046] The anti-neurodegenerative disease or neuroprotective activity of conopeptide JG001 is demonstrated in animal models of Parkinson's disease. Conopeptide JG001 is effective in reversing the behavioral deficits induce by dopamine depletion. Conopeptide JG001 shows behavioral potentiation, especially locomotor activity. Conopeptide JG001 enhances the effect of L-DOPA in reversing the behavioral deficits induce by dopamine depletion. Thus, conopeptide JG001 is effective neuroprotective agents and anti-neurodegenerative disease agents.

[0047] The effect of conopeptide JG001 on muscle control is demonstrated in animals. At low doses, conopeptide JG001 is effective in hampering voiding at the level of the urethra. At higher doses, conopeptide JG001 is effective in eliminating all lower urinary tract activity. In

the animal studies, it appears that conopeptide JG001 is more discriminatory in their inhibitory effects on striated sphincter than on bladder when compared with other NMDA antagonists. Thus, conopeptide peptide JG001 can be dosed in such a way so as to selectively decrease bladder/sphincter dyssynergia, especially in spinal cord injured patients, and are therefore useful
5 for treating urinary incontinence and muscle relaxation.

[0048] In addition to the above medical uses, several of the conopeptides of the present invention have agricultural uses. The conopeptides derived from worm hunting *Conus* species contain N-terminal sequences distinctive from that of piscivorous species in that residue 2 is invariably aromatic. These peptidic toxins are directed at invertebrate glutamate receptors and therefore have agricultural applications, e. for the control of nematodes, parasitic worms and other worms.

[0049] Pharmaceutical compositions containing a compound of the present invention as the active ingredient can be prepared according to conventional pharmaceutical compounding techniques. See, for example, *Remington's Pharmaceutical Sciences*, 18th Ed. (1990, Mack Publishing Co., Easton, PA). Typically, an antagonistic amount of active ingredient will be admixed with a pharmaceutically acceptable carrier. The carrier may take a wide variety of forms depending on the form of preparation desired for administration, e.g., intravenous, oral, parenteral or intrathecally. For examples of delivery methods see U.S. Patent No. 5,844,077, incorporated herein by reference.

20 [0050] "Pharmaceutical composition" means physically discrete coherent portions suitable for medical administration. "Pharmaceutical composition in dosage unit form" means physically discrete coherent units suitable for medical administration, each containing a daily dose or a multiple (up to four times) or a sub-multiple (down to a fortieth) of a daily dose of the active compound in association with a carrier and/or enclosed within an envelope. Whether the
25 composition contains a daily dose, or for example, a half, a third or a quarter of a daily dose, will depend on whether the pharmaceutical composition is to be administered once or, for example, twice, three times or four times a day, respectively.

[0051] The term "salt", as used herein, denotes acidic and/or basic salts, formed with inorganic or organic acids and/or bases, preferably basic salts. While pharmaceutically
30 acceptable salts are preferred, particularly when employing the compounds of the invention as medicaments, other salts find utility, for example, in processing these compounds, or where non-medicament-type uses are contemplated. Salts of these compounds may be prepared by art-recognized techniques.

[0052] Examples of such pharmaceutically acceptable salts include, but are not limited to, inorganic and organic addition salts, such as hydrochloride, sulphates, nitrates or phosphates and acetates, trifluoroacetates, propionates, succinates, benzoates, citrates, tartrates, fumarates, maleates, methane-sulfonates, isothionates, theophylline acetates, salicylates, respectively, or the like. Lower alkyl quaternary ammonium salts and the like are suitable, as well.

[0053] As used herein, the term "pharmaceutically acceptable" carrier means a non-toxic, inert solid, semi-solid liquid filler, diluent, encapsulating material, formulation auxiliary of any type, or simply a sterile aqueous medium, such as saline. Some examples of the materials that can serve as pharmaceutically acceptable carriers are sugars, such as lactose, glucose and sucrose, starches such as corn starch and potato starch, cellulose and its derivatives such as sodium carboxymethyl cellulose, ethyl cellulose and cellulose acetate; powdered tragacanth; malt, gelatin, talc; excipients such as cocoa butter and suppository waxes; oils such as peanut oil, cottonseed oil, safflower oil, sesame oil, olive oil, corn oil and soybean oil; glycols, such as propylene glycol, polyols such as glycerin, sorbitol, mannitol and polyethylene glycol; esters such as ethyl oleate and ethyl laurate, agar; buffering agents such as magnesium hydroxide and aluminum hydroxide; alginic acid; pyrogen-free water; isotonic saline, Ringer's solution; ethyl alcohol and phosphate buffer solutions, as well as other non-toxic compatible substances used in pharmaceutical formulations.

[0054] Wetting agents, emulsifiers and lubricants such as sodium lauryl sulfate and magnesium stearate, as well as coloring agents, releasing agents, coating agents, sweetening, flavoring and perfuming agents, preservatives and antioxidants can also be present in the composition, according to the judgment of the formulator. Examples of pharmaceutically acceptable antioxidants include, but are not limited to, water soluble antioxidants such as ascorbic acid, cysteine hydrochloride, sodium bisulfite, sodium metabisulfite, sodium sulfite, and the like; oil soluble antioxidants, such as ascorbyl palmitate, butylated hydroxyanisole (BHA), butylated hydroxytoluene (BHT), lecithin, propyl gallate, aloha-tocopherol and the like; and the metal chelating agents such as citric acid, ethylenediamine tetraacetic acid (EDTA), sorbitol, tartaric acid, phosphoric acid and the like.

[0055] For oral administration, the compounds can be formulated into solid or liquid preparations such as capsules, pills, tablets, lozenges, melts, powders, suspensions or emulsions. In preparing the compositions in oral dosage form, any of the usual pharmaceutical media may be employed, such as, for example, water, glycols, oils, alcohols, flavoring agents, preservatives, coloring agents, suspending agents, and the like in the case of oral liquid preparations (such as,

for example, suspensions, elixirs and solutions); or carriers such as starches, sugars, diluents, granulating agents, lubricants, binders, disintegrating agents and the like in the case of oral solid preparations (such as, for example, powders, capsules and tablets). Because of their ease in administration, tablets and capsules represent the most advantageous oral dosage unit form, in 5 which case solid pharmaceutical carriers are obviously employed. If desired, tablets may be sugar-coated or enteric-coated by standard techniques. The active agent can be encapsulated to make it stable to passage through the gastrointestinal tract while at the same time allowing for passage across the blood brain barrier. See for example, WO 96/11698.

[0056] For parenteral administration, the compound may be dissolved in a pharmaceutical carrier and administered as either a solution or a suspension. Illustrative of suitable carriers are water, saline, dextrose solutions, fructose solutions, ethanol, or oils of animal, vegetative or synthetic origin. The carrier may also contain other ingredients, for example, preservatives, suspending agents, solubilizing agents, buffers and the like. When the compounds are being administered intrathecally, they may also be dissolved in cerebrospinal fluid.

[0057] A variety of administration routes are available. The particular mode selected will depend of course, upon the particular drug selected, the severity of the disease state being treated and the dosage required for therapeutic efficacy. The methods of this invention, generally speaking, may be practiced using any mode of administration that is medically acceptable, 20 meaning any mode that produces effective levels of the active compounds without causing clinically unacceptable adverse effects. Such modes of administration include oral, rectal, sublingual, topical, nasal, transdermal or parenteral routes. The term "parenteral" includes subcutaneous, intravenous, epidural, irrigation, intramuscular, release pumps, or infusion.

[0058] For example, administration of the active agent according to this invention may 25 be achieved using any suitable delivery means, including:

- (a) pump (see, e.g., Luer & Hatton (1993), Zimm et al. (1984) and Ettinger et al. (1978));
- (b), microencapsulation (see, e.g., U.S. Patent Nos. 4,352,883; 4,353,888; and 5,084,350);
- (c) continuous release polymer implants (see, e.g., U.S. Patent No. 4,883,666);
- (d) macroencapsulation (see, e.g., U.S. Patent Nos. 5,284,761, 5,158,881, 4,976,859 and 4,968,733 and published PCT patent applications WO92/19195, WO 95/05452);
- (e) naked or unencapsulated cell grafts to the CNS (see, e.g., U.S. Patent Nos. 5,082,670 and 5,618,531);

(f) injection, either subcutaneously, intravenously, intra-arterially, intramuscularly, or to other suitable site; or

(g) oral administration, in capsule, liquid, tablet, pill, or prolonged release formulation.

[0059] In one embodiment of this invention, an active agent is delivered directly into the
5 CNS, preferably to the brain ventricles, brain parenchyma, the intrathecal space or other suitable CNS location, most preferably intrathecally.

[0060] Alternatively, targeting therapies may be used to deliver the active agent more specifically to certain types of cell, by the use of targeting systems such as antibodies or cell specific ligands. Targeting may be desirable for a variety of reasons, e.g. if the agent is unacceptably toxic, or if it would otherwise require too high a dosage, or if it would not otherwise be able to enter the target cells.

[0061] The active agents, which are peptides, can also be administered in a cell based delivery system in which a DNA sequence encoding an active agent is introduced into cells designed for implantation in the body of the patient, especially in the spinal cord region. Suitable delivery systems are described in U.S. Patent No. 5,550,050 and published PCT Application Nos. WO 92/19195, WO 94/25503, WO 95/01203, WO 95/05452, WO 96/02286, WO 96/02646, WO 96/40871, WO 96/40959 and WO 97/12635. Suitable DNA sequences can be prepared synthetically for each active agent on the basis of the developed sequences and the known genetic code.

20 [0062] The active agent is preferably administered in an therapeutically effective amount. By a "therapeutically effective amount" or simply "effective amount" of an active compound is meant a sufficient amount of the compound to treat the desired condition at a reasonable benefit/risk ratio applicable to any medical treatment. The actual amount administered, and the rate and time-course of administration, will depend on the nature and
25 severity of the condition being treated. Prescription of treatment, e.g. decisions on dosage, timing, etc., is within the responsibility of general practitioners or specialists, and typically takes account of the disorder to be treated, the condition of the individual patient, the site of delivery, the method of administration and other factors known to practitioners. Examples of techniques and protocols can be found in *Remington's Pharmaceutical Sciences*.

30 [0063] Dosage may be adjusted appropriately to achieve desired drug levels, locally or systemically. Typically the active agents of the present invention exhibit their effect at a dosage range from about 0.001 mg/kg to about 250 mg/kg, preferably from about 0.01 mg/kg to about 100 mg/kg of the active ingredient, more preferably from about 0.05 mg/kg to about 75 mg/kg.

A suitable dose can be administered in multiple sub-doses per day. Typically, a dose or sub-dose may contain from about 0.1 mg to about 500 mg of the active ingredient per unit dosage form. A more preferred dosage will contain from about 0.5 mg to about 100 mg of active ingredient per unit dosage form. Dosages are generally initiated at lower levels and increased 5 until desired effects are achieved. In the event that the response in a subject is insufficient at such doses, even higher doses (or effective higher doses by a different, more localized delivery route) may be employed to the extent that patient tolerance permits. Continuous dosing over, for example 24 hours or multiple doses per day are contemplated to achieve appropriate systemic levels of compounds.

[0064] For the treatment of pain, if the route of administration is directly to the CNS, the dosage contemplated is from about 1 ng to about 100 mg per day, preferably from about 100 ng to about 10 mg per day, more preferably from about 1 μ g to about 100 μ g per day. If administered peripherally, the dosage contemplated is somewhat higher, from about 100 ng to about 1000 mg per day, preferably from about 10 μ g to about 100 mg per day, more preferably from about 100 μ g to about 10 mg per day. If the conopeptide is delivered by continuous infusion (e.g., by pump delivery, biodegradable polymer delivery or cell-based delivery), then a lower dosage is contemplated than for bolus delivery.

[0065] Advantageously, the compositions are formulated as dosage units, each unit being adapted to supply a fixed dose of active ingredients. Tablets, coated tablets, capsules, 20 ampoules and suppositories are examples of dosage forms according to the invention.

[0066] It is only necessary that the active ingredient constitute an effective amount, i.e., such that a suitable effective dosage will be consistent with the dosage form employed in single or multiple unit doses. The exact individual dosages, as well as daily dosages, are determined according to standard medical principles under the direction of a physician or veterinarian for 25 use humans or animals.

[0067] The pharmaceutical compositions will generally contain from about 0.0001 to 99 wt. %, preferably about 0.001 to 50 wt. %, more preferably about 0.01 to 10 wt.% of the active ingredient by weight of the total composition. In addition to the active agent, the pharmaceutical compositions and medicaments can also contain other pharmaceutically active compounds. 30 Examples of other pharmaceutically active compounds include, but are not limited to, analgesic agents, cytokines and therapeutic agents in all of the major areas of clinical medicine. When used with other pharmaceutically active compounds, the conopeptides of the present invention may be delivered in the form of drug cocktails. A cocktail is a mixture of any one of the

compounds useful with this invention with another drug or agent. In this embodiment, a common administration vehicle (e.g., pill, tablet, implant, pump, injectable solution, etc.) would contain both the instant composition in combination supplementary potentiating agent. The individual drugs of the cocktail are each administered in therapeutically effective amounts. A 5 therapeutically effective amount will be determined by the parameters described above; but, in any event, is that amount which establishes a level of the drugs in the area of body where the drugs are required for a period of time which is effective in attaining the desired effects.

10 [0068] The practice of the present invention employs, unless otherwise indicated, conventional techniques of chemistry, molecular biology, microbiology, recombinant DNA, genetics, immunology, cell biology, cell culture and transgenic biology, which are within the skill of the art. See, e.g., Maniatis *et al.*, 1982; Sambrook *et al.*, 1989; Ausubel *et al.*, 1992; Glover, 1985; Anand, 1992; Guthrie and Fink, 1991; Harlow and Lane, 1988; Jakoby and Pastan, 1979; *Nucleic Acid Hybridization* (B. D. Hames & S. J. Higgins eds. 1984); *Transcription And Translation* (B. D. Hames & S. J. Higgins eds. 1984); *Culture Of Animal Cells* (R. I. Freshney, Alan R. Liss, Inc., 1987); *Immobilized Cells And Enzymes* (IRL Press, 1986); B. Perbal, *A Practical Guide To Molecular Cloning* (1984); the treatise, *Methods In Enzymology* (Academic Press, Inc., N.Y.); *Gene Transfer Vectors For Mammalian Cells* (J. H. Miller and M. P. Calos eds., 1987, Cold Spring Harbor Laboratory); *Methods In Enzymology*, 20 Vols. 154 and 155 (Wu *et al.* eds.), *Immunochemical Methods In Cell And Molecular Biology* (Mayer and Walker, eds., Academic Press, London, 1987); *Handbook Of Experimental Immunology*, Volumes I-IV (D. M. Weir and C. C. Blackwell, eds., 1986); Riott, *Essential Immunology*, 6th Edition, Blackwell Scientific Publications, Oxford, 1988; Hogan *et al.*, Manipulating the Mouse Embryo, (Cold Spring Harbor Laboratory Press, Cold Spring Harbor, 25 N.Y., 1986).

EXAMPLES

30 [0069] The present invention is described by reference to the following Examples, which are offered by way of illustration and are not intended to limit the invention in any manner. Standard techniques well known in the art or the techniques specifically described below were utilized.

EXAMPLE 1

Isolation of DNA Encoding Conopeptide JG001

[0070] DNA coding for conopeptide JG001 (Gly-Xaa₁-Asp-Xaa₁-Val-Ser-Gln-Met-Ser-Xaa₂-Xaa₁-Ile-Leu-Arg-Xaa₁-Leu-Glu-Leu-Gln-Xaa₂; Xaa₁ and Xaa₂ are as X₁ and X₂ above; SEQ ID NO:33); was isolated and cloned in accordance with conventional techniques. The DNA was isolated by reverse transcription-PCR using *Conus aurisiacus* venom duct mRNA and primer CCon8 as the forward primer and the primer LibU as the reverse primer. The sequences for these primers are as follows:

CCon8: CAGGATCCTGTATCTGCTGGTGCCCCTGGTG (SEQ ID NO:34) and
 LibU: AAGCTCGAGTAACAAACGCAGAGT (SEQ ID NO:35).

EXAMPLE 2

In vivo Activity of Conopeptide JG001 in Frings Audiogenic Seizure Susceptible Mice

[0071] *In vivo* anticonvulsant activity of conopeptide JG001 (in which Xaa₁ and Xaa₂ are each Gla) was analyzed in Frings audiogenic seizure susceptible mice as described by White et al. (1992). The results for conopeptide JG001 are shown in Tables 1-3.

TABLE 1Effect of Conopeptide JG001 on the Audiogenic Seizure Susceptibility of Frings Mice Following i.c.v. Administration

Dose (pmol, i.c.v.)	# Protected / # Tested		# Toxic / # Tested	
	30 min.	120 min.	30 min.	120 min.
300	4 / 4	4 / 4	0 / 4	0 / 4
1000	3 / 4	4 / 4	2 / 4	1 / 4

Ref: HA2:142-143

TABLE 2Time Effect of Conopeptide JG001 Against Audiogenic Seizure Susceptibility of Frings Mice Following i.c.v. Administration

# Prot. / # Tested	Dose 75 pmol	Time (hrs)					Reference HA2:143
		1 / 4	1 / 2	1	2	4	
	--	4 / 4	--	--	3 / 4	--	
# Toxic / # Tested	75 pmol	--	0 / 4	--	0 / 4	--	HA2:143

TABLE 3

Effect of Conopeptide JG001 on the Audiogenic Seizure
Susceptibility of Frings Mice Following i.c.v. Administration

5	Dose (pmol)	Seizure Score \pm S.E.M.	# Protected / # Tested (at 30 min)	ED ₅₀ (pmol)	# Tested (at 30 min)	TD ₅₀ (pmol)
	18.75	5 \pm 0	0 / 8			
	37.5	3.25 \pm 0.86	3 / 8			
	56.25	2.5 \pm 0.95	4 / 8	46.79		
10	75	0.13 \pm 0.13	8 / 8	(33.82-58.33)*		
	300				1 / 8	
	1000				3 / 8	

* 95% confidence interval

Ref: HA2:142-145

[0072] Conopeptide JG001 yielded an effective dose (ED₅₀) of 46.79 pmol, with a 95% confidence interval of 33.82-58.33 pmol. Furthermore, conopeptide JG001 yielded a toxic dose (TD₅₀) of 1000 pmol (toxicity to 3/8 animals). The dose required to elicit neurotoxicity was >20 times greater than the effective dose (TD₅₀/ED₅₀ = 1000/46.79 = 21.37 = Protective Index, PI). The therapeutic dose of typical anti-seizure medications is close to the toxic dose (typical PI = 2-3). Since the protective index is high for conopeptide JG001, this peptide will be better tolerated than previous anti-convulsant agents.

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EXAMPLE 3

In vivo Activity of Conopeptide JG001 in CF No. 1 Mice

[0073] *In vivo* anticonvulsant activity of conopeptide JG001 is analyzed in CF No. 1 mice as described by White et al. (1995), using the maximal electroshock, subcutaneous pentylenetetrazole (Metrazol) seizure threshold and threshold tonic extension test. Conopeptide JG001 is found to have anticonvulsant activity.

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EXAMPLE 4

In Vivo Activity of Conopeptide JG001 in Pentylenetetrazole-Induced Threshold Seizure Model

[0074] The *in vivo* activity of conopeptide JG001 is analyzed using timed intravenous infusion of pentylenetetrazole (White et al., 1995). At time to peak effect, the convulsant solution (0.5% pentylenetetrazole in 0.9% saline containing 10 U.S.P. units/ml heparin sodium) is

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infused into the tail vein at a constant rate of 0.34 ml/min. The time in seconds from the start of the infusion to the appearance of the first twitch and the onset of clonus is recorded for each drug treated or control animal. The times to each endpoint are converted to mg/kg of pentylenetetrazole for each mouse, and mean and standard error of the mean are calculated. It is 5 found that conopeptide JG001 elevates the i.v. pentylenetetrazole seizure threshold.

EXAMPLE 5

In Vivo Activity of Conopeptide JG001 in Parkinson's Disease Animal Model

10 [0075] The anti-Parkinsonian potential of conopeptide JG001 is examined in rats with unilateral lesions of the nigrostriatal dopamine system. The unilateral lesions are created by local infusion of the neurotoxin 6-hydroxydopamine (6-OHDA) into the right substantia nigra of anesthetized rats. The rats recovered for two weeks at which time they are anesthetized and guide cannulae implanted into the brain, ending in the right lateral ventricle. The guide cannulae are kept patent with a stylet placed in the guide cannula. One week later, the rats are placed in a cylindrical Plexiglas® cage, the stylet is removed, and an infusion cannula is inserted into the guide. The infusion cannula is attached to a syringe on an infusion pump which delivered conopeptide JG001 (0.5 mM, 5.0 mM or 50 mM) or control vehicle at a rate of 1 μ l/min for a total injection of 2 μ l (1 nmol/2 μ l). Fifteen minutes after the injection of conopeptide JG001, L-Dopa (4 mg/kg ip) is injected. The number of full rotations contralateral and ipsilateral to the 20 dopamine-depleted hemisphere is then counted for 2 minutes, every 10 minutes, for 2 hours. A video of the rats is also made to follow the behavioral potentiation of the treatment. It is seen that the tested compound reverses the behavioral deficits induced by dopamine depletion. In addition to the above tests, the *in vivo* activity of conopeptide JG001 in combination with SKF 38393 is compared with that of SKF 38393 alone. It is seen that the combination of conopeptide 25 JG001 and SKF 38393 demonstrates increased activity.

EXAMPLE 6

In vivo Activity of Conopeptide JG001 in Pain Models

30 [0076] The anti-pain activity of conopeptide JG001 is shown in several animal models. These models include the nerve injury model (Chaplan, et al., 1997), the nociceptive response to s.c. formalin injection in rats (Codene, 1993) and an NMDA-induced persistent pain model (Liu, et al., 1997). In each of these models it is seen that the conopeptides and conopeptide derivatives have analgesic properties.

[0077] More specifically, this study evaluates the effect of intrathecal administration of conopeptide JG001 in mice models of nociceptive and neuropathic pain. For nociceptive pain, the effect of the conopeptide JG001 is studied in two different tests of inflammatory pain. The first is the formalin test, ideal because it produces a relatively short-lived, but reliable pain behavior that is readily quantified. There are two phases of pain behavior, the second of which is presumed to result largely from formalin-evoked inflammation of the hind paw. Conopeptide JG001 is administered 10 minutes prior to injection of formalin. The number of flinches and/or the duration of licking produced by the injection is monitored. Since the first phase is presumed to be due to direct activation of primary afferents, and thus less dependent on long term changes in the spinal cord, conopeptide JG001 is presumed to have greatest effect on the magnitude of pain behavior in the second phase.

[0078] The mechanical and thermal thresholds in animals that received an injection of complete Freund's adjuvant into the hind paw are also studied. This produces a localized inflammation including swelling of the hind paw and a profound decrease in mechanical and thermal thresholds, that are detected within 24 hours after injection. The changes in thresholds in rats that receive conopeptide JG001 are compared with those of rats that receive vehicle intrathecal injections.

[0079] To evaluate the contribution of long term, NMDA receptor-mediated changes to neuropathic (i.e., nerve injury-induced) behavior, a modification of the Seltzer model of pain that has been adapted for the mouse is used. A partial transection of the sciatic nerve is first made. This also produces a significant drop in mechanical and thermal thresholds of the partially denervated hind paw. In general, the mechanical changes are more profound. They peak around 3 days after surgery and persist for months.

[0080] An important issue is whether the drugs are effective when administered after the pain model has been established, or whether they are effective only if used as a pretreatment. Clearly, the clinical need is for drugs that are effective after the pain has developed. To address this issue, animals are studied in which conopeptide JG001 is administered repeatedly, after the inflammation (CFA) or nerve injury has been established. In these experiments, conopeptide JG001 is injected daily by the intrathecal (i.t.) route. The mechanical and thermal thresholds (measured, respectively, with von Frey hairs in freely moving animals and with the Hargreave's test, also in freely moving animals) are repeated for a 2 to 4 week period after the injury is induced and the changes in pain measured monitored over time.

EXAMPLE 7

Isolation of DNA Encoding Conopeptides

[0081] DNA coding for conopeptides was isolated and cloned in accordance with conventional techniques using general procedures well known in the art, such as described in Example 1 or in Olivera et al. (1996). Alternatively, cDNA libraries was prepared from *Conus* venom duct using conventional techniques. DNA from single clones was amplified by conventional techniques using primers which correspond approximately to the M13 universal priming site and the M13 reverse universal priming site. Clones having a size of approximately 300-500 nucleotides were sequenced and screened for similarity in sequence to known conopeptides similar to conopeptide JG001 isolated in Example 1. The DNA sequences, encoded propeptide sequences and sequences of the mature toxins are set forth in Table 4. DNA sequences coding for the mature toxin can also be prepared on the basis of the DNA sequences set forth on these pages. An alignment of the conopeptides of the present invention with respect to conantokin G is set forth in Table 5. An alignment of the peptides of the present invention is set forth in Table 6.

TABLE 4

Name: Conotoxin-C1

Species: catus

Cloned: Yes

DNA Sequence:

GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCCACCTA
 20 ATCCTAGGCACGGGCACACTAGATCATGGAGGCGCACTGACTGAACGCCGTTGGG
 TGACGCCACAGCGCTGAGACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
 GCACCGACGACAGTGGCAAGGCACAGGTTGACTCAGATGAAGAGGATTCTCAAAAAA
 GCAAGGAAACACGGCTAAAAGCGACGAAGAGCTACTACGAGAGGATGTAGAGACT
 GTTTAGAACTCGAAAGGAATGGAAAAAGATAATCAAGCTGAGTGTCCACGTGAC
 30 ACTCGTCAGTTCTAAAGTCCCCAGATAATCGTTCCCTATTTGCCACATTCTTCTT
 TCTCTTTCATTTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:36)

Translation:

MQLTYTYLLVPLVTFHLILGTGTLHDGGALTERRSGDATA LRPEPVLLQKSAARSTDD
 35 SGKDRLTQMKRILKKQGNTAKSDEELLREDVETVLELERNGKR (SEQ ID NO:37)

Toxin Sequence:

Ser-Asp-Xaa1-Xaa1-Leu-Leu-Arg-Xaa1-Asp-Val-Xaa1-Thr-Val-Leu-Xaa1-Leu-Xaa1-Arg-
 40 Asn-# (SEQ ID NO:38)

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Name: Conotoxin-C2
Species: catus
Cloned: Yes

5 **DNA Sequence:**

GCGATGCAACTGTACACGTATCTGTATCTGCTGGTGCACCTTCCACCTA
 ATCCTAGGCACGGGCACACTAGATCATGGAGGCGCACTGACTGAACGCCGTTGGG
 TGACGCCACAGCGCTGAGACCTGAGCCTGTCCTCTGCAGAAATCCGCTGCCGCA
 GCACCGACGACAGTGGCAAGGCACAGGTTGACTCAGATGAAGAGGATTCTCAAAAAA
 10 GCAAGGAAACACGGCTAAAGGCACGAAGAGCTACTACGAGAGGATGTAGAGACT
 GTTTAGAACCTGAAAGGGATGGAAAAAGATAATCAAGCTGAGTGTCCACGTGGC
 ACTCGTCAGTTCTAAAGTCCCCAGATAAAATCGTTCCCTATTTGCCACATTCTTCTT
 TCTCTTTCATTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:39)

Translation:

MQLYTYLYLLVPLVTFHLILGTGTLHDGGALTERSGDATA LRPEPVLLQKSAARSTDD
 SGKDRLTQMKRILKKQGNTAKGDEELLREDVETVLELERDGKR (SEQ ID NO:40)

20 **Toxin Sequence:**

Gly-Asp-Xaa1-Xaa1-Leu-Leu-Arg-Xaa1-Asp-Val-Xaa1-Thr-Val-Leu-Xaa1-Leu-Xaa1-Arg-
 Asp-# (SEQ ID NO:41)

25 **Name:** Conotoxin-C3
Species: catus
Cloned: Yes

DNA Sequence:

GCGATGCAACTGTACACGTATCTGTATCTGCTGGCACCTTCCACCTA
 30 ATCCTAGGCACGGGCACACTAGATCATGGAGGCGCACTGACTGAACGCCGTTGGG
 TGACGCCACAGCGCTGAGACCTGAGCCTGTCCTCTGCAGAAATCCGCTGCCGCA
 GCACCGACGACAGTGGCAAGGCACAGGTTGACTCAGATGAAGAGGATTCTCAAAAAA
 GCAAGGAAACACGGCTAAAGGCACGAAGAGCTACTACGAGAGGATGTAGAGACT
 GTTTAGAACCCGAAAGGAATGGAAAAAGATAATCAAGCTGAGTGTCCACGTGAC
 35 ACTCGTCAGTTCTAAAGTCCCCAGATAAAATCGTTCCCTATTTGCCACATTCTTCTT
 TCTCTTTCATTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:42)

Translation:

MQLYTYLYLLAPLVTFHLILGTGTLHDGGALTERSGDATA LRPEPVLLQKSAARSTDD
 40 SGKDRLTQMKRILKKQGNTAKSDEELLREDVETVLEPERNGKR (SEQ ID NO:43)

45 **Toxin Sequence:**

Ser-Asp-Xaa1-Xaa1-Leu-Leu-Arg-Xaa1-Asp-Val-Xaa1-Thr-Val-Leu-Xaa1-Xaa3-Xaa1-Arg-
 Asn-# (SEQ ID NO:44)

Name: Conotoxin-C4
Species: catus

Cloned: Yes

DNA Sequence:

5 GCGATGCAACTGtACACGTATCTGTATCTGCTGGTGTCCCTGGTACCTTCCACCTAA
 TCCTAGGCACGGGCACACTAGATCATGGAGGCGCACTGACTGAACGCCGTTGGCT
 GACGCCACAGCGCTGGAAGCTGAGCCTGTCCTGCAGAAATCCGCTGCCCGCAG
 CACCGACAAACAATGGCAAGGACAGGTCGACTCAGATGAGGAGGATTCTCAAAAAG
 CAAGGAAACACGGCTAGAATCGAGGAAGGTCTGATAGAGGATCTGGAGACCGCTA
 10 GAGAACGCGACAGTGGAAAAAGATAATCAAGCTGAGTGTCCACGTGACACTCATC
 AGTTCTAAAGTCCCCAGATAAAATCGTCCCTATTTGCCACATTCTTCTTCCTCTT
 TTCGTTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:45)

Translation:

15 MQLYTYLYLLVSLVTFHILGTGTLHDGGALTERRLADATALEAEPVLLQKSAARSTD
 NNGKDRSTQMRRILKKQGNTARIEEGLIEDLETARERDSGKR (SEQ ID NO:46)

Toxin Sequence:

20 Ile-Xaa1-Xaa1-Gly-Leu-Ile-Xaa1-Asp-Leu-Xaa1-Thr-Ala-Arg-Xaa1-Arg-Asp-Ser-# (SEQ ID NO:47)

25 **Name:** Conotoxin-C5

Species: catus

Cloned: Yes

DNA Sequence:

30 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTGTCCCTGGTACCTTCCACCTA
 ATCCTAGGCACGGGCACACTAGATCATGGAGGCGCACTGACTGAACGCCGTTGGC
 TGACGCCACAGCGCTGGAAGCTGAGCCTGTCCTGCAGAAATCCGCTGCCCGCA
 GCACCGACAAACAATGGCAAGGACAGGTCGACTCAGATGAGGAGGATTCTCAAAAAG
 GCAAGGAAACACGGCTAGAATCGAGGAAGGTCTGATAGAGGATCTGGAGGCTGCT
 AGAGAACGCGACAGTGGAAAAAGATAATCAAGCTGAGTGTCCACGTGACACTCAT
 CAGTTCTAAAGTCCCCAGATAAAATCGTCCCTATTTGCCACATTCTTCTTCCTCT
 35 TTTCGTTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:48)

Translation:

40 MQLYTYLYLLVSLVTFHILGTGTLHDGGALTERRLADATALEAEPVLLQKSAARSTD
 NNGKDRSTQMRRILKKQGNTARIEEGLIEDLEAARERDSGKR (SEQ ID NO:49)

Toxin Sequence:

45 Ile-Xaa1-Xaa1-Gly-Leu-Ile-Xaa1-Asp-Leu-Xaa1-Ala-Ala-Arg-Xaa1-Arg-Asp-Ser-# (SEQ ID NO:50)

45 **Name:** Conotoxin-C6

Species: catus

Cloned: Yes

DNA Sequence:

5 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTGCCCTGGTGACCTTCCACCTA
 ATCCTAGGCACGGGCACACTAGATCATGGAGGGCGCACTGACTGAACGCCGTTGGC
 TGACGCCACAGCGCTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
 GCACCGACGACAATGGCAAAGACAGGTTGACTCACATGAAGAGGATTCTCAAAAAA
 ACGAGCAAACAAAGCCAGAGGCGAACAGAGTTGAAGCATAACCGGAGGCAGTA
 AGACAAACAAGAATGTATAAGAAATAATAATCGACCTTGGTGTCCAAGTGACA
 CTCGTCAGTTCTAAAGTCTCCAGATAGATCGTCCCTATTTGCCACACTCTTCTT
 TCTCTTCATTTAAGTCCCCAAATCTTCATGTTATT (SEQ ID NO:51)

10 **Translation:**
 MQLYTYLYLLVPLVTFHLILGTGTLHGGALTERRSADATAALKPEPVLLQKSAARSTDD
 NGKDRLTHMKRJLKRRANKARGEPEVGSIPEAVRQQECIRNNNNRPWCPK (SEQ ID
 NO:52)

15 **Toxin Sequence:**
 Gly-Xaa1-Xaa3-Xaa1-Val-Gly-Ser-Ile-Xaa3-Xaa1-Ala-Val-Arg-Gln-Gln-Xaa1-Cys-Ile-Arg-
 Asn-Asn-Asn-Asn-Arg-Xaa3-Xaa4-Cys-Xaa3-Lys-^ (SEQ ID NO:53)

20 **Name:** Conotoxin-Bu1
Species: bullatus
Cloned: Yes

25 **DNA Sequence:**
 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTGCCCTGGTGACCTTCCACCTA
 ATCCTGGGCACGGGCACACTAGATCATGGAGGGCGCACTGACTGAACGCCGTTGGC
 TGACGCCACAGCACTGAAACCTGAGCCTGTCCTCCTGCAGAAAACCGCTGCCGCA
 GCACCGACGACAATGGCAAGAAGAGGCTGACTCAGAGGAAGAGGATTCTCAAAAAA
 GCGAGGAAACACGGCTAGAAACCCGAAACTTATATAGAGATTGGAGATTCTA
 30 GGGAACTCGAAGAGATTGGAAAAAAGATAATCAAGCTGGTGTCCACGTGACACTC
 GTCAGTTCTGAAGTCCCGAGGTAGATCGTCCCTATTTGCCACACTCTTCTTCT
 CTTTCATTTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:54)

35 **Translation:**
 MQLYTYLYLLVPLVTFHLILGTGTLHGGALTERRSADATAALKPEPVLLQKTAARSTDD
 NGKKRLTQRKRJLKRRGNTARNPETYIEIVEISRELEEEIGKR (SEQ ID NO:55)

40 **Toxin Sequence:**
 Asn-Xaa3-Xaa1-Thr-Xaa5-Ile-Xaa1-Ile-Val-Xaa1-Ile-Ser-Arg-Xaa1-Leu-Xaa1-Xaa1-Ile-#
 (SEQ ID NO:56)

45 **Name:** Conotoxin-Bu2
Species: bullatus
Cloned: Yes

DNA Sequence:
 GCGATGCAACTGTACACGTATCTGTATTGCTGGTGCCCTGGTGACCTTCCACCTA
 ATCCTGGGCACGGGCACACTAGATCATGGAGGGCGCACTGACTGAACGCCGTTGGC

5 TGACGCCACAGCGCTGAAACCTGAGCCTGTCCTCCTGCAGAAAACCGCTGCCGCA
 GCACCGACGACAATGGCAAGAAGAGGCTGACTCAGAGGAAGAGGATTCTCAAAAAA
 GCGAGGAAACACGGCTAGAAACCCCGAAACTTATTATAATTAGAGCTTGTGGAGA
 TTTCTAGGAACTCGAAGAAATTGGAAAAAGATAATCAAGCTGGGTGTTCCACGTG
 10 ACACTCGTCAGTTCTAAGTCCCAGGGTAGATCGTTCCCTATTTTGCACACTCTT
 CTTTCTCTTCATTAATTCCCCAAACTTCATGTTATT (SEQ ID NO:57)

Translation:

10 MQLTYTYLLVPLVTFHLILGTGTLHGGALTERSADATALKPEPVLLQKTAARSTDD
 NGKKRLTQRKRILKKRGNTARNPETYYNLELVEISRELEEGKR (SEQ ID NO:58)

Toxin Sequence:

Asn-Xaa3-Xaa1-Thr-Xaa5-Xaa5-Asn-Leu-Xaa1-Leu-Val-Xaa1-Ile-Ser-Arg-Xaa1-Leu-Xaa1-Xaa1-Ile-# (SEQ ID NO:59)

15 **Name:** Conotoxin-Bt1
Species: betulinus
Cloned: Yes

20 **DNA Sequence:**

25 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTGCCCTGGTGACCTTCTACCTA
 ATCCTAGGCACGGGCACGCTAGGTCATGGAGGGCGCACTGACTGAACGCCGTTGGC
 TGATGCCACAGCGCTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCCGCCGCA
 GCACCGACGACAATGGCAAGGACAGGTTGACTCAGATGATCAGGATTCTCAAAAAAG
 CGAGGAAACATGGCCAGAGGCGGCGAAGAAGTTAGAGAGTCTGCAGAGACTCTTC
 ATGAACTCACGCCGTAGGAAAAAGAAAAAGATTAATCAAGCTGGGTGTTCCACGTG
 ACACTCGTCAGTTCTAAAGTCCCCAGTTCCATCTTGCCACGTTCTTTCTTTTC
 ATTCAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:60)

30 **Translation:**

35 MQLTYTYLLVPLVTFYLILGTGTLHGGALTERLADATALKPEPVLLQKSAARSTDD
 NGKDRLTQMIRILKKRGNMARGGEVRESAETLHELTP (SEQ ID NO:61)

Toxin Sequence:

40 Gly-Gly-Xaa1-Xaa1-Val-Arg-Xaa1-Ser-Ala-Xaa1-Thr-Leu-His-Xaa1-Leu-Thr-Xaa3-^ (SEQ ID NO:62)

45 **Name:** Conotoxin-Bt2
Species: betulinus
Cloned: Yes

DNA Sequence:

45 GCGATGCAACTGTACGTATCTGTATCTGCTGGTGCCGCTGGTGACCTTCTACCTA
 ATCCTAGGCACGGGCACGCTAGGTCATGGAGGGCGCACTGACTGAACGCCGTTGGC
 TGACGCCACAGCGCTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCCGCCGCA
 GCACTGACGACAATGGCAAGGACAGGTTGACTCAGATGATCAGGATTCTCAAAAAAG
 CGAGGAAACATGGCCAGAGGCGGCGAAGAAGTTAGAGAGTCTGCAGAGACTCTTC

ATGAAATCACGCCGTAGGAAAAAGAAAAAGATTAATCAAGCTGGGTGTTCCACGTG
 ACACTCGCCAGTTCTAAAGTCCCCAGTTCCATCTTGCCAGGTTCTTCTCTTT
 CATTCAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:63)

5 **Translation:**

MQLYTYLYLLVPLVTFYLILGTGTLGHGGALTERRLADATALKPEPVLLQKSAARSTDD
 NGKDRLTQMIRILKKRGNMARGGEVRESAETLHEITP (SEQ ID NO:64)

Toxin Sequence:

10 Gly-Gly-Xaa1-Xaa1-Val-Arg-Xaa1-Ser-Ala-Xaa1-Thr-Leu-His-Xaa1-Ile-Thr-Xaa3-^ (SEQ ID NO:65)

Name: Conotoxin-Bt3
Species: betulinus
Cloned: Yes

DNA Sequence:

20 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCTACCTA
 ATCCTAGGCACGGGCACGCTAGGTATGGAGGCGCACTGACTGAACGCCGTTGGC
 TGACGCCACAGCGCTGAAACCTAACGCCTATCCTCCTGCAGAAATCCGCCGCCGCA
 GCACTGACGACAATGGCAAGGACAGGTTGACTCAGATGATCAGGATTCTAAAAAG
 CGAGGAAACATGGGCAGAGACGGCGAAGAAGTCAGAGAGGCTGCAGAGACTCTTA
 ATGAACTCAGCCGTAGGAAAAAGAAAAAGATTAATCAAGCTGGGTGTTCCACGTG
 ACACTCGTCAGTTCTAAAGTACCCAGTTCCATCTTGCCACGTTCTTTCTTT
 CATTCAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:66)

Translation:

MQLYTYLYLLVPLVTFYLILGTGTLGHGGALTERRLADATALKPKPILLQKSAARSTDD
 30 NGKDRLTQMIRILKKRGNMGRDGEVREAAETLNELTP (SEQ ID NO:67)

Toxin Sequence:

35 Asp-Gly-Xaa1-Xaa1-Val-Arg-Xaa1-Ala-Ala-Xaa1-Thr-Leu-Asn-Xaa1-Leu-Thr-Xaa3-^ (SEQ ID NO:68)

Name: Conotoxin-Bt4
Species: betulinus
Cloned: Yes

DNA Sequence:

40 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCCACCTA
 ATCCTAGGCACGGGCACGCTAGGTATGGAGGCGCACTGACTGAAGGCCGTTGGC
 TGACGCCACAGCACTGAAACCAGGGCCTGCCTCCTGCAGAAATCCGCTGCCGCA
 GCACCGACGACAATGGCAAGGACAGGTTGACTCAGATGAAGAGGACTCTAAAAAA
 GCGAGGAAACACGGCCAGAGGCTACGAAGATGATAGAGAGATTGCAGAGACTGTT
 AGAGAACTCGAGGAAGCAGGAAAATGAAAAAGATTAATCAAGCTGGGTGTTCCAC
 GTGACACTTGTCAAGTTCTAAAGTCCCCAGATAGATCGTCCCTATTTTGCCACATT
 TTTTTCTCTTTCTTTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:69)

Translation:

MQLYTYLYLLVPLVTFHLILGTGTLGHGGALTESRSADATALKPGPVLLQKSAARSTDD
NGKDRLTQMCKRTLKKRGNTARGYEDDREIAETVRELEEAAGK (SEQ ID NO:70)

5

Toxin Sequence:

Gly-Xaa5-Xaa1-Asp-Asp-Arg-Xaa1-Ile-Ala-Xaa1-Thr-Val-Arg-Xaa1-Leu-Xaa1-Xaa1-Ala-#
(SEQ ID NO:71)

10

Name: Conotoxin-Bt5
Species: betulinus
Cloned: Yes

DNA Sequence:

GCGATGCAACTGTACACGTATCTGTATCTGCTGGTGCCGCTGGTACCTTCTACCTA
ATCCTAGGCACGGGCACGCTAGGTCATGGAGGGCGCACTGACTGAACGCCGTTGGC
TGACGCCACAGCGCTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCCGCCGCA
GCACTGACGACAATGGCAAGGACAGGTTGACTCAGATGATCAGGATTCTCAAAAAG
CGAGGAAACATGGCCAGAGGCAGGGAGAAGTTAGAGAGTCTGCAGAGACTCTC
ATGAAATCACGCCGTAGGAAAAAGAAAAAGATTAATCAAGCTGGGTGTTCCACGTG
ACACTCGTCAGTTCTAAAGTCCCCAGTTCCATCTTGCCAGGTTCTTCTCTTT
CATTCAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:72)

25

Translation:

MQLYTYLYLLVPLVTFYLILGTGTLGHGGALTERRLADATALKPEPVLLQKSAARSTDD
NGKDRLTQMIRILKKRGNMARGGGEVRESAETLHEITP (SEQ ID NO:73)

Toxin Sequence:

30

Gly-Gly-Gly-Xaa1-Val-Arg-Xaa1-Ser-Ala-Xaa1-Thr-Leu-His-Xaa1-Ile-Thr-Xaa3-^ (SEQ ID NO:74)

Name: Conotoxin-Af6
Species: ammiralis
Cloned: Yes

DNA Sequence:

35

GCGATGCAACTGTACACGTATCTGTGTCTGCTGGTGCCCTGGTACCTTCTACCTA
ATTCTAGGCACGGGCACACTAGCTCATGGAGGGCGCACTGACCGAACGCCGTTGGC
TCACGCCAGAGTAATAGAACCTGATCCTGCCCTGGAGAAACTCCGCTCTCCGCA
GCATCCGACGACAACGACAAGGACAGGATGACTCAGAGGAAGAGGATTCTCAAAA
AGTGATGAAACACGGCCAGAGGCAGGGCGAAAGAAGATAGAAATAATGCGGAGGCTGT
TAGAGAAAGACTCGAAGAAATAGGAAAAAGGTAATCAAGCTGGGTGTTCACGTG
40 45
ACACTCATCAGTTCTAAAGTCCCCAGATAGATCGTTCCATTTGCCATATTCTT
CCTTCTCTTCATGTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:75)

Translation:

MQLYTYLCLLVPLVTFYLILGTGTLAHHGALTERRLAHARVIEPDPAPLENSALRSIRRQ
RQGQDDSEEEDSQKVMKHGQRRERR (SEQ ID NO:76)

Toxin Sequence:

5 Xaa2-Gly-Gln-Asp-Asp-Ser-Xaa1-Xaa1-Xaa1-Asp-Ser-Gln-Lys-Val-Met-Lys-His-Gly-Gln-
Arg-Arg-Xaa1-Arg-Arg-^ (SEQ ID NO:77)

10 Name: Conotoxin-Ep1

Species: episcopatus

Cloned: Yes

DNA Sequence:

15 GCGATGCAACTGTACACGTATCTGTGCTGGTCCCCCTGGTGACCTTCTACCTA
ATTCTAGGCACGGGCACACTAGCTCATGGAGGCGCACTGACTGAACATCGTTCGGC
CGACGCCACAGCACTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
GCACCGACGACAACGGCAAGGACAGGTTGACTCGGTGGAAGGGGATTCTCAAAAAA
GCGAGGAAACACGGCCAGAGGCCGGAAAGATATTGTGGAGACTATTACAGAACTC
GAAAAAAATAGGAAAAAGGTAATCAAGCTGGGTGTTCCACGTGACACTCATCAGTTC
20 TAAAGTCCCCAGATAGATCGTCCCTATTTGCCATTCTTCTTTCTTTATG
TAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:78)

Translation:

25 MQLYTYLCLLVPLVTFYLILGTGTLAHHGALTEHRSADATALKPEPVLLQKSAARSTDD
NGKDRLTRWKGILKKRGNNTARGGKDIVETITELEKIGKR (SEQ ID NO:79)

Toxin Sequence:

30 Gly-Gly-Lys-Asp-Ile-Val-Xaa1-Thr-Ile-Thr-Xaa1-Leu-Xaa1-Lys-Ile-# (SEQ ID NO:80)

30 Name: Conotoxin-L1

Species: lynceus

Cloned: Yes

DNA Sequence:

35 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCCACCTA
ATCCTAGGCACGGGCACACTAGATCATGGAGGCGCACTGACTGAACGCCGTTCGAC
TGATGCCATAGCACTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCTCTGCCCGCAG
40 CACCGACGATAATGGCAACGACAGGTTGACTCAGATGAAGAGGATCCTCAAAAAG
CGAGGAAACAAAGCCAGAGGCCGAAGAAGAAGTTGCAAAATGGCGGCAGAGATTG
CCAGAGAAAACGCTGCAAATGGGAAATGATAATCAAGTTGGGTGTTCCACGTGACA
CTCGTCAGTTCTAAAGTCCCCAGATAGATCGTCCCTATTTGCCACATTCTTCTT
TCTCTTTCATTTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:81)

Translation:

45 MQLYTYLYLLVPLVTFHLILGTGTLHGGALTERRSTDIAALKPEPVLLQKSSARSTDD
NGNDRLTQMKRILKKRGNKARGEVEVAKMAAEIARENAANGK (SEQ ID NO:82)

Toxin Sequence:

Gly-Xaa1-Xaa1-Xaa1-Val-Ala-Lys-Met-Ala-Ala-Xaa1-Ile-Ala-Arg-Xaa1-Asn-Ala-Ala-Asn-#
(SEQ ID NO:83)

5 **Name:** Conotoxin-L2
Species: lynceus
Cloned: Yes

DNA Sequence:

10 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCTACCTA
ATCCTAGGCACGGGCACGCTAGGTATGGAGGCACACTGACTGAACGCCGTTCGGC
TGATGCCACAGCACTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
GCACCGGCACGATGCCAAGGAGAGGTTGACTCAGACGAAGAGGATTGCAAAAAA
GCGAGCAAACACGACCAGAGGCAAAGAAGAGGATAGAGAGAGATTGAGACTGTT
AGAGAACTCGAAGAAATAGGAAAAAGATGATCAAGCTGGGTGTTCCACGTGACAC
TCGTCAGTTCAAAGTCCCCAGATAGATCGTTCCCTATTTGCCACATTCTTCTTT
CTTTTTCATTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:84)

Translation:

20 MQLYTYLYLLVPLVIFYLILGTGTLGHGGLTERRSADATALKPEPVLLQKSAARSTGD
DAKERLTQTKRIRKKRANTTRGKEEDREIVETVRELEEIFKR (SEQ ID NO:85)

Toxin Sequence:

25 Gly-Lys-Xaa1-Xaa1-Asp-Arg-Xaa1-Ile-Val-Xaa1-Thr-Val-Arg-Xaa1-Leu-Xaa1-Xaa1-Ile-#
(SEQ ID NO:86)

30 **Name:** Conotoxin-L3
Species: lynceus
Cloned: Yes

DNA Sequence:

35 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCCACCTA
ATCCTAGGCACGGGCACACTAGATCATGGAGGCACACTGACTGAACGCCGTTCGAC
TGACGCCATAGCACTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCTCTGCCGCA
CACCAGACAATGGCAACGACAGGTTGATTCAAGATGAAGAGGATTCTCAAAAAGC
GAGGAAACAAAGCCAGAGGCGAAGAGGAAGTTGCAAAATGGCGGCAGAGCTTAC
CAGAGAAGAAGCTGTAAGGGAAATGATAATCAAGTTGGGTGTTCCACGTGACAC
TCGTCAGTTCAAAGTCCCCAGATAGATCGTTCCCTATTTGCCACATTCTTCTTT
40 CTATTTCATTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:87)

Translation:

45 MQLYTYLYLLVPLVTFHLILGTGTLDHGGALTERRSTDIAALKPEPVLLQKSSARSTDD
NGNDRLIQMKRILKKRGNKARGEVEAKMAELTREEAVKGK SEQ ID NO:88)

Toxin Sequence:

46 Gly-Xaa1-Xaa1-Xaa1-Val-Ala-Lys-Met-Ala-Ala-Xaa1-Leu-Thr-Arg-Xaa1-Xaa1-Ala-Val-Lys-#
(SEQ ID NO:89)

5 **Name:** Conotoxin-Fi1

Species: *figulinus*

Cloned: Yes

DNA Sequence:

10 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTACCTTCTACCTA
 ATCCTAGGCACGGGCACGCTAGGTCATGGAGGCGCACTGACTGAACGCCGTTGGC
 TGACGCCACAGCGCTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
 GCACCGACGACAATGACAAGGACAGGCTGACCCAGATGAAGAGGATTTCAAAAAA
 GCGAGGAAACAAAGCCAGAGGCAGGAAGAAGTTGCAGAGATGGCGGCAGAGATT
 GCAAGAGAAAATCAAGCAAACGGGAAAAGATAATCAAACGGGTGTTCCACGTGA
 CACTCGTCAGTTCTAAAGTCCCCAGATAGGTCGTTCTATGTTGCCACATTCTTC
 TTTTCTTTCATTAATTCCCCAAATCTTCATGTTTATT (SEQ ID NO:90)

Translation:

15 MQLYTYLYLLVPLVTFYLILGTGLHGGALTERLADATALKPEPVLLQKSAARSTD
 NDKDRLTQMKRIFKKRGNKARGEVEAEMAAEIARENQANGKR (SEQ ID NO:91)

20 **Toxin Sequence:**

Gly-Xaa1-Xaa1-Xaa1-Val-Ala-Xaa1-Met-Ala-Ala-Xaa1-Ile-Ala-Arg-Xaa1-Asn-Gln-Ala-Asn-#
 (SEQ ID NO:92)

25 **Name:** Conotoxin-Fi2

Species: *figulinus*

Cloned: Yes

30 **DNA Sequence:**

ATCCTAGGCACGGGCACACTAGCTCATGGAGGCGCACCGACTGAACGCCGTTGGC
 TGACACCCACAGCACTGAAACCCGAGCATGTCCTCCTGCAGATGTCGCTGCCGCA
 GCACCAACGATAATGGCAAGGACAGGTTGACTCAGATGAAGAGGATTCTCAAAAAAA
 GCAAGGAAACACAGCCAGAAGCTACGAACAAGCTAGAGAAGTTCAGGAGGCTGTT
 35 AATGAACACTCAAGGAAAGAGGTAAAAAGATAATCATGCTGGGTGTTCCACGTGACAC
 TCGTCAGTTCTAAAGCCCCCAGATAGATTGTTCCGTATTTTACACGTTCTTCTTT
 CTCTTTCATTAATTCCCCAAATCTTCATGTTTATT (SEQ ID NO:93)

40 **Translation:**

MQLYTYLYLLVPLVTFYLILGTGLAHGGAPTERLADTTALKPEHVLLQMSAARSTN
 DNGKDRLTQMKRILKKQGNTARSYEQAREVQEAVNELKERGKKIIMLGVPDRTRQF
 (SEQ ID NO:94)

45 **Toxin Sequence:**

Ser-Xaa5-Xaa1-Gln-Ala-Arg-Xaa1-Val-Gln-Xaa1-Ala-Val-Asn-Xaa1-Leu-Lys-Xaa1-Arg-#
 (SEQ ID NO:95)

Name: Conotoxin-Fi2a
Species: *figulinus*
Cloned: Yes

5 DNA Sequence:

GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTACCTTCTACCTA
 ATCCTAGGGACGGGCACACTAGCTCATGGAGGCGCACCGACTGAACGCCGTTGGC
 TGACACCACAGCACTGAAACCCGAGCATGTCCTCCTGCAGATGTCCGCTGCCGCA
 GCACCAACGATAATGGCAAGGCAGAGCTACGAACAAGCTAGAGAAAGTTCAAGGAGGCTGTT
 10 GCAAGGAAACACAGCCAGAAGCTACGAACAAGCTAGAGAAAGTTCAAGGAGGCTGTT
 AATGAACTCAAGGAAAGAGGTAAAAAGATAATCATGCTGGGTGTTCCACGTGACAC
 TCGTCAGTTCTAAAGCCCCAGATAGATTGTTCCGTATTTTACACGTTCTTCTTT
 CTCTTTCATTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:96)

Translation:

MQLTYLYLLVPLVTYIILGTGTLAHGGAPERRLADTTALKPEHVLLQMSAARSTN
 DNGKDRLTQMKRILKKQGNTARSYEQAREVQEAVNELKERGKKIIMLGVPRDTRQF
 (SEQ ID NO:97)

20 Toxin Sequence:

Ser-Xaa5-Xaa1-Gln-Ala-Arg-Xaa1-Val-Gln-Xaa1-Ala-Val-Asn-Xaa1-Leu-Lys-Xaa1-Arg-Gly-
 Lys-Lys-Ile-Ile-Met-Leu-Gly-Val-Xaa3-Arg-Asp-Thr-Arg-Gln-Phe-^ (SEQ ID NO:98)

25 Name: Conotoxin-Fi3
Species: *figulinus*
Cloned: Yes

DNA Sequence:

GCGATGCAACTGTACACGTATCTGTATCTGCTGGtGCCCTGGTACGTTCCACCTA
 30 ATCCTAGGCACGGGCACACTAGCTCATGGAGGCGCACTGGCTGAACGCCGTTGGC
 TGACGCCACAGCGCTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
 GCACCGACGACAATGGCAAGGACAGGTTGACTGAGATGAAGAGGATTCTCAAAAAA
 GCGAGGAAACACGCCAGAGACTACGAAGATGATAGAGAGATTGCAGAGACTGTT
 35 AGAGAACTCGAAGAAATAGTAAAAGATAATCAAGCTGGGTGTTCAATTGACACTC
 ATCAGTTCTAAAGTCCCCAGATAGATCGTTCCCTAATTGCCCACGTTCTTCTTTCT
 CTTTCATTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:99)

Translation:

MQLTYLYLLVPLVTFHILGTGTLAHGGALAERRLADATALKPEPVLLQKSAARSTD
 DNGKDRLTEMKRILKKRGNTARDYEDDREIAETVRELEEIGKR (SEQ ID NO:100)

40 Toxin Sequence:

Asp-Xaa5-Xaa1-Asp-Asp-Arg-Xaa1-Ile-Ala-Xaa1-Thr-Val-Arg-Xaa1-Leu-Xaa1-Xaa1-Ile-#
 45 (SEQ ID NO:101)

Name: Conotoxin-Fi4
Species: *figulinus*

Cloned: Yes

DNA Sequence:

5 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTGCCTCTGGTACCTTCTACCTA
 ATCCTAGGCACGGGCACGCTAGGTATGGAGGGCGCACTGACTGAACGCCGTTGGC
 TGACGCCACAGCGCTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
 GCACCGACGACAATGGCAAGGACAGGTTGACTCAGATGAAGGGGACTGTAAAAAA
 GCGAGGAAACACGGCCGAAGAAGTTAGAGAGGGCTGCAGAGACTCTTCATGAACCT
 CGCTGTAGGAAAAAGAAAAAGATTAATCAAGCTGGGTGTTCCACGTGACACTCGTC
 10 AGTTCTAAAGTCCCCAGTTCCCTATCTTGCCACGTTTCTTCTTTCATCCAA
 TTCCCCAAATCTTCATGTTATT (SEQ ID NO:102)

Translation:

MQLYTYLYLLVPLVTFYLILGTGTLGHGGALTERRLADATAALKPEPVLLQKSAARSTD
 NGKDRLTQMKGTVKRGNTAEEVREAAETLHESL (SEQ ID NO:103)

Toxin Sequence:

Gly-Asn-Thr-Ala-Xaa1-Xaa1-Val-Arg-Xaa1-Ala-Ala-Xaa1-Thr-Leu-His-Xaa1-Leu-Ser-Leu-[^]
 (SEQ ID NO:104)

Name: Conotoxin-Fi5

Species: *figulinus*

Cloned: Yes

DNA Sequence:

5 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTGCCTCTGGTACCTTCCACCTA
 ATCCTAGGCACGGGCACACTAGGTATGGAGGGCGCACTGACTGAACGCCGTTGGC
 TGACGCCACAGCGCTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
 30 GCACCGACGTCAATGGCAAGGACAGGTTGACTGAGATGAAGAGGATTCTAAAAAA
 GCGAGGAAGCATATCCATGGGCTTCGAACATAGAAGAGAGATTGCAGAGTTGGTTA
 GAGAACTCGCTGAAATAGGTAAACGATAATCAAGCTGGGTGTTCCACTAACACTCG
 TCAGTTCTAAAGTCCCCAGATAGATCGTCCCTATCTTGCCACATTTCCTCTT
 TTCATTAAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:105)

35

Translation:

MQLYTYLYLLVPLVTFHILGTGTLGHGGALTERRLADATAALKPEPVLLQKSAARSTD
 NGKDRLTEMKRILKKRGSISMGEHRREIAELVRELAEGKR (SEQ ID NO:106)

40

Toxin Sequence:

Gly-Ser-Ile-Ser-Met-Gly-Phe-Xaa1-His-Arg-Arg-Xaa1-Ile-Ala-Xaa1-Leu-Val-Arg-Xaa1-Leu-
 Ala-Xaa1-Ile-# (SEQ ID NO:107)

45

Name: Conotoxin-Di1

Species: *distans*

Cloned: Yes

DNA Sequence:

5 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGGCCTTGCACCTA
 ATCCAAGGCACGGGCACACTAGGCCATGGAGGCGCACTGACTGAAGGCCGTTCGGC
 TGACGCCACAGCGCCGAAACCTGAACCTGTCTCCTGCAGAAAATCCGATGCCGCA
 GCGCCGACGACAACCGCAAGGACAAGTTGACTCAGATGAAGAGGACTCTGAAAAAA
 GCAAGGACACATTGCCAGAACCATAACTGCTGAAGAGGCAGAGAGGACTAGTGAA
 AGAATGTCATCAATGGAAAAAGATAATCAAGCTGGGTGTTCCACGTGACACTCGT
 CAGTTCTAAAGTCCCCAGATAAAATCGTCCCTGTTTGCCTGTTCTTCTCTTCTT
 TTCATTCAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:108)

10 **Translation:**

MQLYTYLYLLVPLVAFHLIQGTGTLGHGGALTEGRSADATAPKPEPVLLQKSDARSAD
 DNGKDKLTQMKRTLKKQGHIARTITAEERTSERMSSMGKR (SEQ ID NO:109)

15 **Toxin Sequence:**

Thr-Ile-Thr-Ala-Xaa1-Xaa1-Ala-Xaa1-Arg-Thr-Ser-Xaa1-Arg-Met-Ser-Ser-Met-# (SEQ ID NO:110)

20 **Name:** Conotoxin-Di2

Species: distans

Cloned: Yes

25 **DNA Sequence:**

30 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTATCCCTGGTGGCCTTGCACCTA
 ATCCAAGGAACGGGCACGCTAGGCCATGGAGGCGCACTGACTGAAGGCCGTTCGGC
 TGACGCCACAGCGCCGAAACCTGAACCTGTCTCGTGCAGAAAATCGGATGCCGCA
 GCGCCGACGACAACCGCAAGGACAAGTTGACTCAGATGAAGAGGATTCTGAAAAAA
 GCAAGAAACCCCAACTCCTGAAGAGGTAGAGCGCCATACCGAAAGACTCAAAAGC
 ATGGGAAAAAGATAATCAAGCTGGGTGTTCCACGTGACACTCGTCAGTTCTAAAGT
 CCCCAGATGGATCGTCCCTGTTTGCCTGTTCTTCATTCAATTCAATT
 CCCAAATCTTCATGTTATT (SEQ ID NO:111)

35 **Translation:**

MQLYTYLYLLVSLVAFHLIQGTGTLGHGGALTEGRSADATAPKPEPVLVQKSDARSAD
 DNRKDKLTQMKRILKKQETPTPEEVERHTERLKSMGKR (SEQ ID NO:112)

40 **Toxin Sequence:**

Xaa2-Xaa1-Thr-Xaa3-Thr-Xaa3-Xaa1-Xaa1-Val-Xaa1-Arg-His-Thr-Xaa1-Arg-Leu-Lys-Ser-
 Met-# (SEQ ID NO:113)

45 **Name:** Conotoxin-P1

Species: purpurascens

Cloned: Yes

DNA Sequence:

GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTACCTTGCACCTA
 ATCCTAGGCACGGGAATGCTAGCTCATGGAGACACACTGACTGAACGCCGTTCGGT
 TGACGCCACAGCACTGAAACCTGAGCCTGTCCTGCAGAAAATCCGCTGCCGCA

5 GCACCGACGACAATGACAAGGACAGGTTGACTCAGATGAAGAGGATTCTAAAAAA
 GCGAGGAAACAAAGCCAGAGGCGAAGAAGAACATTCCAAGTATCAAGAGTGTCTT
 AGAGAAGTAAGAGTAAATAAGGTACAACAAGAATGTTAATCAAGCTGGGTGTTCCA
 CGTGACACTCGTCAGTTCTAAAGTCCCCAGATAGATCGTTCCCGATTTCGCCACAT
 TCTTCTTCTCTTTCATTTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:114)

10 **Translation:**
 MQLYTYLYLLVPLVTFHILGTGMLAHGDLTERRSVDATAALKPEPVLLQKSAARSTD
 DNDKDRLTQMKRILKKRGNKARGEHHSKYQECLREVRVNKVQQEC (SEQ ID NO:115)

15 **Toxin Sequence:**
 Gly-Xaa1-Xaa1-Xaa1-His-Ser-Lys-Xaa5-Gln-Xaa1-Cys-Leu-Arg-Xaa1-Val-Arg-Val-Asn-Lys-
 Val-Gln-Gln-Xaa1-Cys-^ (SEQ ID NO:116)

20 **Name:** Conotoxin-P2
Species: purpurascens
Cloned: Yes

25 **DNA Sequence:**
 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCCACCTA
 ATCCTAGGCACGGGCACACTAGCTCATGGAGGCGCACTGACTGAACGCCGTTCCAC
 TGACGCCACAGCACTGAAACCTGAGCCTGTCCTGCAGGAATCTGATGCCCGCAGCA
 CCGACGACAATGACAAGGACAGGTTGACTCAGATGAAGAGGATTCTAAAAAGCG
 AGGAAACAAAGCCAGAGGCGAAGAAGAACATTCCAAGTATCAGGAGTGTCTAGA
 GAAGTAAGAGTAAATAACGTACAACAAGAATGTTAATCAAGCTGGGTGTTCCACGT
 GACACTCGTCAGTTCTAAAGTCCCCAGATAGATCGTTCCCTATTTTGCACATTCTT
 TCTTCTCTTTCATTTAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:117)

30 **Translation:**
 MQLYTYLYLLVPLVTFHILGTGTLAHHGALTERGSTDATAALKPEPVLQESDARSTDDN
 DKDRLTQMKRILKKRGNKARGEHHSKYQECLREVRVNNVQQEC (SEQ ID NO:118)

35 **Toxin Sequence:**
 Gly-Xaa1-Xaa1-Xaa1-His-Ser-Lys-Xaa5-Gln-Xaa1-Cys-Leu-Arg-Xaa1-Val-Arg-Val-Asn-Asn-
 Val-Gln-Gln-Xaa1-Cys-^ (SEQ ID NO:119)

40 **Name:** Conotoxin-P3
Species: purpurascens
Cloned: Yes

45 **DNA Sequence:**
 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCCACCTA
 ATCCTAAGGCACGGGCACACTAGCTCATGGAGGCGACACTGACTGAACGCCGTTCGAC
 TGACACCCACAGCACTGAAACCTGAGCCTGTCCTGCAGAAATCTGATGCCCGCA
 GCACCGACGACAATGACAAGGACAGGTTGACTCAGATGAAGAGGATTCTAAAAAA
 GCGAGGAAACAAAGCCAGAGGCGAAGAAGAACATTCCAAGTATCAGGAGTGTCTT

AGAGAAATAAGAGTAAATAAGGTACAACAAGAATGTTAATCAAGCTGGGTGTTCCA
 CGTGACACCCGTCAGTTCTAAAGTCCCCAGATAGATCGTTCCCTATTTGCCACAT
 TCTTCTTCTCTTCAATTAAATTCCCCAAATCTTCATGTTTATT (SEQ ID NO:120)

5 **Translation:**

MQLYTYLYLLVPLVTFHLILSTGTLAHGGTLTERRSTDALKPEPVLLQKSDARSTDD
 NDKDRLTQMKRILKKRGNKARGEHHSKYQECLREIRVNKVQQEC (SEQ ID NO:121)

Toxin Sequence:

10 Gly-Xaa1-Xaa1-Xaa1-His-Ser-Lys-Xaa5-Gln-Xaa1-Cys-Leu-Arg-Xaa1-Ile-Arg-Val-Asn-Lys-
 Val-Gln-Gln-Xaa1-Cys-^ (SEQ ID NO:122)

Name: Conotoxin-P4
Species: purpurascens
Cloned: Yes

DNA Sequence:

20 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCCACCTA
 ATCCTAACGACGGGACACTAGCTCATGGAGACACACTGACTGAACGCCGTTCGGT
 TGACGCCACAGCACTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
 GCACCGACGACGATGACAAGGACAGGTTGACTCAGAGGAAGAGGATTCTCAAAAAA
 GCAAGGAAACAAAGCCAGAGGCGAAGCAGAACATTACGCGTTCAGGAGTGTCTT
 AGAGAAATAATGTAATAAGGTACAACAAGAATGTTAATCAAGCTGGGTGTTCTA
 CGTGACACTCGTCAGTTCTAAAGTCCCCAGATAGATCGTTCCCTATTTGCCACAT
 TCTTCTTCTCTTCAATTAAATTCCCCAAATCTTCATGTTTATT (SEQ ID NO:123)

Translation:

30 MQLYTYLYLLVPLVTFHLILSTGTLAHGDTLTERRSDATAALKPEPVLLQKSAARSTDD
 DDKDRLTQQRKRLKKQGNKARGEAEHYAFQECLREINVNVKQQEC (SEQ ID NO:124)

Toxin Sequence:

35 Gly-Xaa1-Ala-Xaa1-His-Xaa5-Ala-Phe-Gln-Xaa1-Cys-Leu-Arg-Xaa1-Ile-Asn-Val-Asn-Lys-
 Val-Gln-Gln-Xaa1-Cys-^ (SEQ ID NO:125)

Name: Conotoxin-P5
Species: purpurascens
Cloned: Yes

DNA Sequence:

40 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTCCCCCTGGTGACCTTCCACCTA
 ATCCTAGGCACGGGAATGCTAGCTCATGGAGACACACTGACTGAACGCCGTTCGGT
 TGACGCCACAGCACTGAAACCTGAGCCTGTCCTCCTGCAGAAATCCGCTGCCGCA
 GCACCGACGCCATGGCAAGGACAGGTTGACTCAGAGGAAGAGGATTCTCAAAAAA
 GCGAGGAAACATGGCCAGGGCTAGAAGAAGATATAGAGTTATTGAGACGATC
 GAAGAAATTGGAAAAAGATAACCAAGCTGGGTGTTCCACGTGACACTCGTCGGTTC

TAAAGTCCCCAGATAGATCGTCACTATTTGCCACATTCTTCTTTCTTTCAATT
TAATTCCCCAAATCTTCATGTTATT (SEQ ID NO:126)

Translation:

5 MQLYTYLYLLVPLVTFHLILGTGMLAHGDTLTERRSVDATALKPEPVLLQKSAARSTD
ANGKDRLTQRKRILKKRGNMARGLEEDIEFIEEIGKR (SEQ ID NO:127)

Toxin Sequence:

Gly-Leu-Xaa1-Xaa1-Asp-Ile-Xaa1-Phe-Ile-Xaa1-Thr-Ile-Xaa1-Ile-# (SEQ ID NO:128)

10

Name: Conotoxin-Sm1
Species: *stercusmuscarum*
Cloned: Yes

DNA Sequence:

15 GCGATGCAACTGTACACGTATCTGTATCTGCTGGTGCCCTGGTGACCTTCCACCTA
ATCCTGGGCACGGGCACACTAGATCATGGAGGGCGCACTGACTGAACGCCGTCGGC
TGACGCCACAGCGCTGAAACCTGAGCCTGTCCTGCAGAAATCCGCTGCCGGCAGCA
20 CCGACGACAACGGCAAGGACAGGTTGACTCAGATGAAGAGGATTCTCAAAAAGCG
AGGAAACACGGCTAGAACATCACCAGAAACTGATATAGAGCTTGTATGAAATTAGAAG
AAATTGGAAAAAGATAATCAAGCTGGGTGTTCCACGTGACACTCGTCAGTTCTGAA
25 GTCCCGAGGTAGATCGTCCCTATTTGCCACATTCTTCTTCTTTCATGTAATT
TCCCCAAATCTTCATGTTATT (SEQ ID NO:129)

Translation:

MQLYTYLYLLVPLVTFHLILGTGTLHGGALTERRSADATALKPEPVLQKSAAGSTDD
NGKDRLTQMKRILKKRGNTRITETDIELVMKLEEIGKR (SEQ ID NO:130)

30

Toxin Sequence:

Ile-Thr-Xaa1-Thr-Asp-Ile-Xaa1-Leu-Val-Met-Lys-Leu-Xaa1-Xaa1-Ile-# (SEQ ID NO:131)

35 Where:

Xaa1 = Glu or γ -Carboxy Glu

Xaa2 = Gln or pyroglu

Xaa3 = Pro or Hydroxy Pro

Xaa4 = Trp (D or L) or Bromo Trp (D or L)

40

Xaa5 = Tyr, ^{125}I -Tyr, Mono-Iodo Tyr, Di-Iodo Tyr, O-sulpho-Tyr or O-Phospho-Tyr

\wedge = Free-carboxyl C-term or Amidated C-term, preferably Free-carboxyl

= Free-carboxyl C-term or Amidated C-term, preferably Amidated

? = Status of C-term not known.

TABLE 5

Alignment of Linear γ -Carboxyglutamate Rich Conotoxins (SEQ ID NO:) With Respect to Conantokin G

	Conantokin G	-----GEXXL-QX-NQXLIRX-KSN# (SEQ ID NO:132)
5	Conotoxin-Af6	ZGQDDSEXXDSQKVMKHGQRERR^ (SEQ ID NO:133)
	Conotoxin-Bt1	-----GGXXV-RX-SAXTLHXLTP^ (SEQ ID NO:134)
	Conotoxin-Bt2	-----GGXXV-RX-SAXTLHXITP^ (SEQ ID NO:135)
	Conotoxin-Bt3	-----DGXXV-RX-AAXTLNXLTP^ (SEQ ID NO:136)
10	Conotoxin-Bt4	-----GY-XDDRX-IAXTVRXLEEA# (SEQ ID NO:137)
	Conotoxin-Bt5	-----GGGXV-RX-SAXTLHXITP^ (SEQ ID NO:138)
	Conotoxin-Bu1	-----NP-XTYIX-IVXISRXLLEI# (SEQ ID NO:139)
	Conotoxin-Bu2	-----NP-XTYY--NLX-LVXISRELEEI# (SEQ ID
	NO:140)	
15	Conotoxin-C1	-----SDXXLLRX-DVXTVLXLERN# (SEQ ID NO:141)
	Conotoxin-C2	-----GDXXXLLRX-DVXTVLXLERD# (SEQ ID NO:142)
	Conotoxin-C3	-----SDXXLLRX-DVXTVLXLXPERN# (SEQ ID NO:143)
	Conotoxin-C4	-----IE-XGLIX-DLXTARXRDS# (SEQ ID NO:144)
	Conotoxin-C5	-----IE-XGLIX-DLXAARXRDS# (SEQ ID NO:145)
20	Conotoxin-C6	-----GEPXVGS--IPXAVRQQECIRNNNNRPWCPK^ (SEQ
	ID NO:146)	
	Conotoxin-D11	-T--ITAXXA--XRTSXRMSM# (SEQ ID NO:147)
	Conotoxin-D12	ZET-PTPXXV--XRHTXRLKSM# (SEQ ID NO:148)
	Conotoxin-Ep1	G--GKDIVXTITX--LXKI# (SEQ ID NO:149)
25	Conotoxin-F11	-----GEXXV-AXMAAXIARXNQAN# (SEQ ID NO:150)
	Conotoxin-F12	-----S-YXQARX-VQXAVNXLKER# (SEQ ID NO:151)
	Conotoxin-F12a	-----S-YXQARX-VQXAVNXLKERGKKIIMLGVPRDTRQF^
	(SEQ ID NO:152)	
	Conotoxin-F13	-----D-YXDDRX-IAXTVRXLEEI# (SEQ ID NO:153)
30	Conotoxin-F14	GNTA--XXV-RX-AAXTLHELS-L^ (SEQ ID NO:154)
	Conotoxin-F15	GSISMG-FXHRRX-IAXLVRELAEI# (SEQ ID NO:155)
	Conotoxin-L1	-----GEXXVAK-MAAXIARXNAAN# (SEQ ID NO:156)
	Conotoxin-L2	-----GKXXD-RX-IVXTVRXLLEI# (SEQ ID NO:157)
	Conotoxin-L3	-----GEXXVAK-MAAXLTRXEAVK# (SEQ ID NO:158)
35	Conotoxin-P1	-----GEXXHSK--YQXCLRXVRVNKVQQEC^ (SEQ ID
	NO:159)	
	Conotoxin-P2	-----GEXXHSK--YQXCLRXVRVNNVQQEC^ (SEQ ID
	NO:160)	
	Conotoxin-P3	-----GEXXHSK--YQXCLRXIRVNKVQQEC^ (SEQ ID
40	NO:161)	
	Conotoxin-P4	-----GEAXHYA--FQXCLRXINVNKVQQEC^ (SEQ ID
	NO:162)	
	Conotoxin-P5	-----GLXXD-IX-FIX-TIXEI# (SEQ ID NO:163)
	Conotoxin-Sm1	-----IT-XTDIXLVMKL--XEI# (SEQ ID NO:164)
45		

X is Glu or Gla, preferably Gla particularly with respect to positions 3 and 4 in conantokin G

TABLE 34

Alignment of Linear γ -Carboxyglutamate Rich Conotoxins¹

5	Conotoxin-Af6	ZGQDDSEEEEDSQKVMKHGQRERRR^ (SEQ ID NO:165)
	Conotoxin-Bt1	G---G----EEVRESAETLHELT-P^ (SEQ ID NO:166)
	Conotoxin-Bt2	G---G----EEVRESAETLHEIT-P^ (SEQ ID NO:167)
	Conotoxin-Bt3	D---G----EEVREAAETLNELT-P^ (SEQ ID NO:168)
	Conotoxin-Bt4	G-----YEDDREIAETVRELEEA# (SEQ ID NO:169)
	Conotoxin-Bt5	G---G----GEVRESAETLHEIT-P^ (SEQ ID NO:170)
10	Conotoxin-Bu1	NPETY-----IEIVEISRELEEI# (SEQ ID NO:171)
	Conotoxin-Bu2	NPETY-----YNLELVEISRELEEI# (SEQ ID NO:172)
	Conotoxin-C1	SDEEL-----LREDVETVLELERN# (SEQ ID NO:173)
15	Conotoxin-C2	GDEEL-----LREDVETVLELERD# (SEQ ID NO:174)
	Conotoxin-C3	SDEEL-----LREDVETVLEPERN# (SEQ ID NO:175)
	Conotoxin-C4	IEEGL-----I-EDLETARERD-S# (SEQ ID NO:176)
20	Conotoxin-C5	IEEGL-----I-EDLEAARERD-S# (SEQ ID NO:177)
	Conotoxin-C6	GEPEVGSIPPEAVRQQECIRNNNNRPWCPK^ (SEQ ID NO:178)
	Conotoxin-Di1	--T--ITAEAAERTSERMSSM# (SEQ ID NO:179)
25	Conotoxin-Di2	ZET--PTPEEVERHTERLKSM# (SEQ ID NO:180)
	Conotoxin-Ep1	G---G-----KDIVETITELEKI# (SEQ ID NO:181)
	Conotoxin-Fi1	GEEEVAE-----MAAEIARENQAN# (SEQ ID NO:182)
	Conotoxin-Fi2	-----S-YEQAREVQEAVNELKER# (SEQ ID NO:183)
	Conotoxin-Fi2a	-----S-YXQAREVQEAVNELKERGKKIIMLGVPDRTRQF^
30	Conotoxin-Fi3	D-----YEDDREIAETVRELEEI# (SEQ ID NO:185)
	Conotoxin-Fi4	GNTA----EEVREAAETLHELS-L^ (SEQ ID NO:186)
	Conotoxin-Fi5	GSISMG-FEHRREIAELVRELAEI# (SEQ ID NO:187)
	Conotoxin-L1	GEEEVAK-----MAAEIARENAAAN# (SEQ ID NO:188)
	Conotoxin-L2	G-----KEEDREIVETVRELEEI# (SEQ ID NO:189)
35	Conotoxin-L3	GEEEVAK-----MAAELTREEAVK# (SEQ ID NO:190)
	Conotoxin-P1	GEEEHSKYQECLREVRVNKVQQEC^ (SEQ ID NO:191)
	Conotoxin-P2	GEEEHSKYQECLREVRVNNVQQEC^ (SEQ ID NO:192)
	Conotoxin-P3	GEEEHSKYQECLREIRVNKVQQEC^ (SEQ ID NO:193)
40	Conotoxin-P4	GEAEHYAFQECLREINVNKVQQEC^ (SEQ ID NO:194)
	Conotoxin-P5	G---LEEDIEFIETIE-----EI# (SEQ ID NO:195)
45	Conotoxin-Sm1	-----ITETDIELVMKL---EEI# (SEQ ID NO:196)

¹ The sequences are compared prior to γ -carboxylation.

[0082] It will be appreciated that the methods and compositions of the instant invention
50 can be incorporated in the form of a variety of embodiments, only a few of which are disclosed

herein. It will be apparent to the artisan that other embodiments exist and do not depart from the spirit of the invention. Thus, the described embodiments are illustrative and should not be construed as restrictive.

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